



# Edexcel GCSE (9-1) **Chemistry**

Mark Levesley   Iain Brand   Nigel Saunders   Sue Robilliard   John Ling

# Contents

Teaching and learning

iv

<b>SC1</b>	<b>States of Matter</b>	
<b>SC2</b>	<b>Methods of Separating and Purifying Substances (Paper 1)</b>	<b>1</b>
<b>SC1a</b>	States of matter	2
<b>SC2a</b>	Mixtures	4
<b>SC2b</b>	Filtration and crystallisation	6
<b>SC2c</b>	Paper chromatography	8
<b>SC2d</b>	Distillation	10
<b>SC2d</b>	Core practical – Investigating inks	12
<b>SC2e</b>	Drinking water	14
<b>SC1–2</b>	Preparing for your exams	16

<b>SC3</b>	<b>Atomic Structure (Paper 1 and Paper 2)</b>	<b>17</b>
<b>SC3a</b>	Structure of an atom	18
<b>SC3b</b>	Atomic number and mass number	20
<b>SC3c</b>	Isotopes	22
<b>SC3</b>	Preparing for your exams	24

<b>SC4</b>	<b>The Periodic Table (Paper 1 and Paper 2)</b>	<b>25</b>
<b>SC4a</b>	Elements and the periodic table	26
<b>SC4b</b>	Atomic number and the periodic table	28
<b>SC4c</b>	Electronic configurations and the periodic table	30
<b>SC4</b>	Preparing for your exams	32

<b>SC5</b>	<b>Ionic Bonding</b>	
<b>SC6</b>	<b>Covalent Bonding</b>	
<b>SC7</b>	<b>Types of Substance (Paper 1 and Paper 2)</b>	<b>33</b>
<b>SC5a</b>	Ionic bonds	34
<b>SC5b</b>	Ionic lattices	36
<b>SC5c</b>	Properties of ionic compounds	38
<b>SC6a</b>	Covalent bonds	40
<b>SC7a</b>	Molecular compounds	42
<b>SC7b</b>	Allotropes of carbon	44
<b>SC7c</b>	Properties of metals	46
<b>SC7d</b>	Bonding models	48
<b>SC5–7</b>	Preparing for your exams	50

<b>SC8</b>	<b>Acids and Alkalis (Paper 1)</b>	<b>51</b>
<b>SC8a</b>	Acids, alkalis and indicators	52
<b>SC8b</b>	Looking at acids	54
<b>SC8c</b>	Bases and salts	56
<b>SC8c</b>	Core practical – Preparing copper sulfate	58
<b>SC8d</b>	Alkalis and balancing equations	60
<b>SC8d</b>	Core practical – Investigating neutralisation	62
<b>SC8e</b>	Alkalis and neutralisation	64
<b>SC8f</b>	Reactions of acids with metals and carbonates	66
<b>SC8g</b>	Solubility	68
<b>SC8</b>	Preparing for your exams	70

<b>SC9</b>	<b>Calculations Involving Masses (Paper 1 and Paper 2)</b>	<b>71</b>
<b>SC9a</b>	Masses and empirical formulae	72
<b>SC9b</b>	Conservation of mass	74
<b>SC9c</b>	Moles	76
<b>SC9</b>	Preparing for your exams	78

<b>SC10</b>	<b>Electrolytic Processes</b>	
<b>SC11</b>	<b>Obtaining and Using Metals</b>	
<b>SC12</b>	<b>Reversible Reactions and Equilibria</b>	
<b>SC13</b>	<b>Transition Metals, Alloys and Corrosion (Paper 1)</b>	<b>79</b>
<b>SC10a</b>	Electrolysis	80
<b>SC10a</b>	Core practical – Electrolysis of copper sulfate solution	82
<b>SC10b</b>	Products from electrolysis	84
<b>SC11a</b>	Reactivity	86
<b>SC11b</b>	Ores	88
<b>SC11c</b>	Oxidation and reduction	90
<b>SC11d</b>	Life cycle assessment and recycling	92
<b>SC12a</b>	Dynamic equilibrium	94
<b>SC13a</b>	Transition metals	96
<b>SC13b</b>	Corrosion	98
<b>SC13c</b>	Electroplating	100
<b>SC13d</b>	Alloying	102
<b>SC13e</b>	Uses of metals and their alloys	104
<b>SC10–13</b>	Preparing for your exams	106



<b>SC14</b>	<b>Quantitative Analysis</b>	
<b>SC15</b>	<b>Dynamic Equilibria, Calculations Involving Volumes of Gases</b>	
<b>SC16</b>	<b>Chemical Cells and Fuel Cells (Paper 1)</b>	107
<b>SC14a</b>	Yields	108
<b>SC14b</b>	Atom economy	110
<b>SC14c</b>	Concentrations	112
<b>SC14d</b>	Titrations and calculations	114
<b>SC14d</b>	Core practical – Acid–alkali titration	116
<b>SC14e</b>	Molar volume of gases	118
<b>SC15a</b>	Fertilisers and the Haber process	120
<b>SC15b</b>	Factors affecting equilibrium	122
<b>SC16a</b>	Chemical cells and fuel cells	124
<b>SC14–16</b>	Preparing for your exams	126

<b>SC22</b>	<b>Hydrocarbons</b>	
<b>SC23</b>	<b>Alcohols and Carboxylic Acids</b>	
<b>SC24</b>	<b>Polymers (Paper 2)</b>	171
<b>SC22a</b>	Alkanes and alkenes	172
<b>SC22b</b>	Reactions of alkanes and alkenes	174
<b>SC23a</b>	Ethanol production	176
<b>SC23b</b>	Alcohols	178
<b>SC23b</b>	Core practical – The combustion of alcohols	180
<b>SC23c</b>	Carboxylic acids	182
<b>SC24a</b>	Addition polymerisation	184
<b>SC24b</b>	Polymer properties and uses	186
<b>SC24c</b>	Condensation polymerisation	188
<b>SC24d</b>	Problems with polymers	190
<b>SC22–24</b>	Preparing for your exams	192

<b>SC17</b>	<b>Groups in the Periodic Table</b>	
<b>SC18</b>	<b>Rates of Reaction</b>	
<b>SC19</b>	<b>Heat Energy Changes in Chemical Reactions (Paper 2)</b>	127
<b>SC17a</b>	Group 1	128
<b>SC17b</b>	Group 7	130
<b>SC17c</b>	Halogen reactivity	132
<b>SC17d</b>	Group 0	134
<b>SC18a</b>	Rates of reaction	136
<b>SC18b</b>	Factors affecting reaction rates	138
<b>SC18b</b>	Core practical – Investigating reaction rates	140
<b>SC18c</b>	Catalysts and activation energy	142
<b>SC19a</b>	Exothermic and endothermic reactions	144
<b>SC19b</b>	Energy changes in reactions	146
<b>SC17–19</b>	Preparing for your exams	148

<b>SC25</b>	<b>Qualitative Analysis: Tests for Ions</b>	
<b>SC26</b>	<b>Bulk and Surface Properties of Matter Including Nanoparticles (Paper 2)</b>	193
<b>SC25a</b>	Flame tests and photometry	194
<b>SC25b</b>	Tests for positive ions	196
<b>SC25c</b>	Tests for negative ions	198
<b>SC25c</b>	Core practical – Identifying ions	200
<b>SC26a</b>	Choosing materials	202
<b>SC26b</b>	Composite materials	204
<b>SC26c</b>	Nanoparticles	206
<b>SC25–26</b>	Preparing for your exams	208
	Glossary	209
	Index	214
	The periodic table of the elements	216

<b>SC20</b>	<b>Fuels</b>	
<b>SC21</b>	<b>Earth and Atmospheric Science (Paper 2)</b>	149
<b>SC20a</b>	Hydrocarbons in crude oil and natural gas	150
<b>SC20b</b>	Fractional distillation of crude oil	152
<b>SC20c</b>	The alkane homologous series	154
<b>SC20d</b>	Complete and incomplete combustion	156
<b>SC20e</b>	Combustible fuels and pollution	158
<b>SC20f</b>	Breaking down hydrocarbons	160
<b>SC21a</b>	The early atmosphere	162
<b>SC21b</b>	The changing atmosphere	164
<b>SC21c</b>	The atmosphere today	166
<b>SC21d</b>	Climate change	168
<b>SC20–21</b>	Preparing for your exams	170

# Teaching and learning

The **topic reference** tells you which part of the course you are in. 'SC17c' means, 'Separate Science, Chemistry, unit 17, topic c'.

The **specification reference** allows you to cross reference against the specification criteria so you know which parts you are covering. References that end in C, e.g. C9.2C, are in Chemistry only, the rest are also in the Combined Science specification criteria.

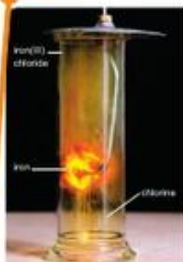
If you see an **H** icon that means that content will be assessed on the Higher Tier paper only.

## SC17c Halogen reactivity

Specification reference: C9.2A; C6.11H; C9.12; C9.13

### Progression questions

- How can displacement reactions be used to work out the reactivity of halogens?
- How can we explain the reactivity of halogens?
- What happens to halogen atoms and halide ions during displacement?



**C** Chlorine displaces iodine from potassium iodide solution.

Table A shows the order of reactivity of different halogens when heated with iron wool. In general the halogens become less reactive as you go down the group.

Halogen	Effect on iron wool
fluorine	burns into flames
chlorine	glows brightly
bromine	glows dull red
iodine	changes colour

**A** halogens and heated iron wool

- Astatine, symbol At, is found below iodine in the halogen group.
  - Write a word equation for the reaction of astatine with iron wool.
  - How will this reaction compare with chlorine? Explain your answer.
- Write a balanced equation for the formation of magnesium iodide (MgI<sub>2</sub>) from its elements.

The order of reactivity of halogens can also be worked out using **displacement reactions**. In a displacement reaction, a more reactive element takes the place of a less reactive element in an ionic compound. So, a more reactive halogen displaces a less reactive halogen from a halide compound. For example, chlorine displaces bromine from sodium bromide in solution, but bromine cannot displace chlorine from sodium chloride.

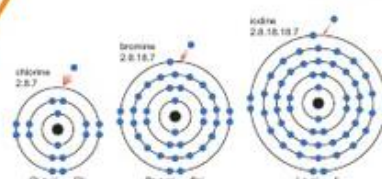


- Which of the following pairs of substances do not react? Explain your choice.
  - $\text{Br}_2 + \text{LiCl}$
  - $\text{Cl}_2 + \text{NaI}$
- Write a balanced equation, with state symbols, for the reaction that occurs in photo C.
- What would be observed if bromine gas were bubbled through potassium iodide solution?

132

To explain the trend in reactivity we need to look at the electronic configuration of the halogen atoms.

Group 7 atoms gain one electron when they react. Down the group, the distance between the outermost shell containing electrons and the nucleus increases. This means that the force of attraction between the positive nucleus and an incoming negative electron decreases, and so the ions do not form so easily and the reactivity decreases.



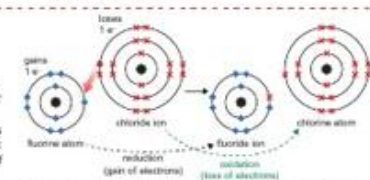
**D** Going down group 7, the outermost electron shell gets further from the nucleus and the ions are less readily formed.

- 6** Explain why fluorine is the most reactive halogen and astatine is the least reactive halogen.

### Redox

When a metal reacts with oxygen it loses electrons and so we can define **oxidation** as a 'loss of electrons'. **Reduction** is the opposite and is a 'gain of electrons'. Use the mnemonic 'OILRIG' to remember that 'Oxidation Is Loss, Reduction Is Gain'. These two processes occur at the same time in displacement reactions, which makes them examples of **reduction-oxidation or redox reactions**.

Diagram E shows the reaction between fluorine atoms and chloride ions. As the fluorine atoms are **reduced**, by gaining electrons, the chloride ions are **oxidised**, by losing electrons.



**E** Fluorine gains electrons more readily than chlorine. It displaces chlorine from chloride ions:  $\text{F}_2(\text{g}) + 2\text{Cl}^-(\text{aq}) \rightarrow 2\text{F}^-(\text{aq}) + \text{Cl}_2(\text{g})$

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Design a summary table or diagram to describe and explain the trend in reactivity of the halogens.

### Extend

**E1** Use a displacement reaction to explain what is meant by 'redox reactions'.

133

By the end of the topic you should be able to confidently answer the **Progression questions**. Try to answer them before you start and make a note of your answers. Think about what you know already and what more you need to learn.

Each question has been given a **Pearson Step** from 1 to 12. This tells you how difficult the question is. The higher the step the more challenging the question.

When you've worked through the main student book questions, answer the **Progression questions** again and review your own progress. Decide if you need to reinforce your own learning by answering the **Strengthen question**, or apply, analyse and evaluate your learning in new contexts by tackling the **Extend question**.



# Paper 1

## SC1 States of Matter / SC2 Methods of Separating and Purifying Substances

Millions of tonnes of tiny bits of plastic are floating in the oceans, and they harm wildlife. Water currents cause the plastic to collect in certain areas. The biggest of these is the 'Great Pacific Garbage Patch' in the Pacific Ocean, which could be three times the area of the UK. At the age of 19, Dutch student Boyan Slat came up with the idea of using giant floating booms to direct the plastic pieces into a mechanism that would filter the plastic out of the water. The idea relies on two properties of the plastic – it floats and it is insoluble in water. Not everyone agrees that it will work, and think that the system would not survive in the oceans. In this unit you will learn about how materials can be separated from one another using their properties.

### The learning journey

Previously you will have learnt at KS3:

- how particles are arranged in solids, liquids and gases and how their energy changes with changes of state
- how mixtures differ from pure substances
- how to separate some mixtures using filtration, distillation and chromatography.

In this unit you will learn:

- how to use information to predict the state of a substance
- how the arrangement, movement and energy of particles change during changes of state
- how to use melting points to tell the difference between mixtures and pure substances
- how to identify substances using melting points and chromatography
- how different methods of separation work
- how to choose a separation method based on the properties of the substances in a mixture.



# SC1a States of matter

Specification reference: C2.1; C2.2; C2.3; C2.4

## Progression questions

- What are particles like in substances in the solid, liquid and gas states?
- What changes happen to particles during the different changes of state?
- How do you decide what state a substance will be in at a given temperature?



**A** This 'ice hotel' is made entirely from ice and snow – these are both water in the solid state.

The three **states of matter** are solid, liquid and gas. For example, water can exist in the solid state as ice, or in the familiar liquid state, or in the gas state as steam or water vapour.

## The particle model

Some **particles** are large enough to see, like the dust on a computer screen. Others, like **atoms** and **molecules**, are far too small for you to see. When chemists discuss particles, they usually mean these very small particles.

The **particle model** explains state changes in a substance in terms of the arrangement, movement and energy stored in its particles.

## Did you know?

Science recognises 16 different types of ice, depending on the arrangement of the water molecules. A type called amorphous ice is found in space (such as on comets). A type called Ice IV is what you'll find in a kitchen freezer, here on Earth.



- 1** Describe the difference in the arrangements of particles in a solid and in a liquid.



- 2** Describe the difference in the movement of particles in a liquid and in a gas.

State	Particle diagram	Arrangement of particles	Movement of particles
Gas		random far apart	fast in all directions
Liquid		random close together	move around each other
Solid		regular close together	vibrate about fixed positions

**B** Particles in the solid state contain the smallest amount of stored energy; particles in the gas state contain the most.



## State changes

State changes are **physical changes**. They can be reversed, and the **chemical properties** of the substance do not change. This is because the particles themselves do not change – only their arrangement, movement and amount of stored energy.

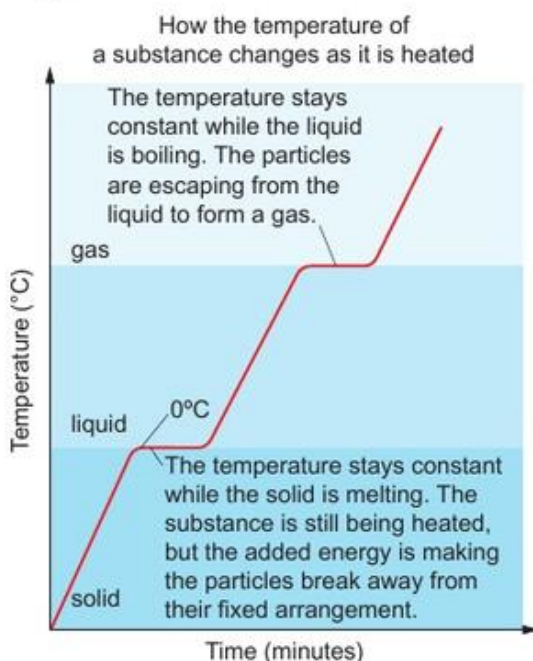
- 2nd** 3 State the meaning of the terms 'sublimation' and 'deposition'.

Particles are attracted to one another by weak forces of attraction. There are many of these forces in a solid. Some of these are overcome during melting. The remaining **attractive forces** between particles in a liquid are overcome during evaporation and boiling (when a substance is evaporating as fast as it can). For this to happen, energy must be transferred from the surroundings to the particles. This is why you heat ice to melt it, and why you boil water in a kettle. Diagram D shows how the temperature changes when water in the solid state is heated until it reaches the gas state.

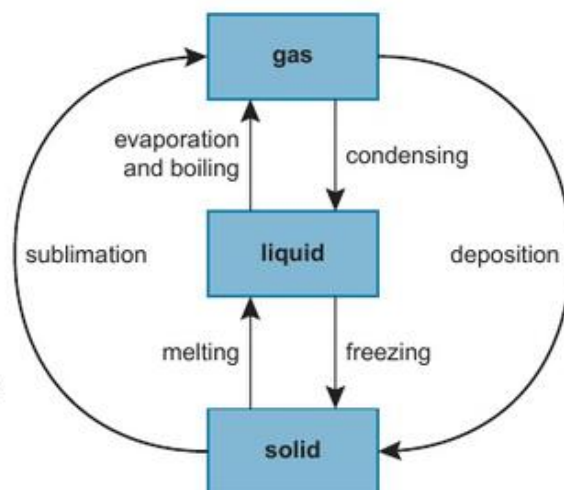
Some attractive forces form between particles during condensing, and many attractive forces are formed during freezing. For this to happen, energy must be transferred from the particles to the surroundings. This is why water vapour turns into water droplets on a cold window, and why you put water in a freezer to make ice.

You can predict the state of a substance if you know its temperature, and its **melting point** and **boiling point**. If the temperature is:

- below the melting point, the substance is solid
- between the melting point and boiling point, the substance is liquid
- above the boiling point, the substance is gas.



D a heating curve for water



C the interconversions between the three states of matter

- 5th** 4 Describe how you can see from a 'heating curve' (such as diagram D) that a substance is changing state.
- 7th** 5 Explain what happens to the particles when a substance melts.
- 6th** 6 The melting point of gallium is  $29.8^{\circ}\text{C}$  and its boiling point is  $2204^{\circ}\text{C}$ . Predict its state at  $25^{\circ}\text{C}$ ,  $100^{\circ}\text{C}$  and at  $2205^{\circ}\text{C}$ .

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Draw a diagram to show the states of matter. On your diagram, name each state change and describe what happens to the particles as it happens.

### Extend

**E1** Explain why the arrangement, movement and energy of particles change during changes of state.

### Exam-style question

Camping gas is used by campers and hikers. It is a mixture of propane and butane. Explain, in terms of the arrangement of fuel particles, why camping gas is stored in cylinders as a liquid rather than as a gas. (2 marks)

# SC2a Mixtures

Specification reference: C2.5; C2.6

## Progression questions

- What is the difference between a pure substance and a mixture?
- What happens to its particles when a solid melts?
- How do melting points allow you to spot the differences between pure substances and mixtures?



**A** You can tell this gold bar is very nearly pure because of the '999.9' stamped on it. A number lower than 1000 on this 'fineness' scale means it is impure.

The composition (make-up) of a **pure** substance:

- cannot be changed
- is the same in all parts of a piece of the substance.

So, for example, pure gold contains only gold atoms.



**1** Which type of atoms are found in a piece of pure silver?



**2** State what is meant by the term 'impure'. Explain what the term 'impure' means.

Gold is an **element** and can be pure, but **compounds** can also be pure. The sugar we use at home is a compound called sucrose. It contains carbon, hydrogen and oxygen atoms chemically bonded together to form sucrose molecules. You cannot change the composition of pure sucrose.

## Did you know?

The purest gold ever was produced in 1957 and was 999.999 on the fineness scale.

Gold purity is still often measured on the older carat scale, where 24 carat gold is pure gold.



**B** Pure sucrose is always sucrose, no matter how finely it is ground down.

A pure substance has the same fixed composition in all its parts and so we can't separate it into other substances using physical methods (such as filtering or picking bits out).

A **mixture** contains elements and/or compounds that are not chemically joined together. You *can* use physical processes to separate mixtures into different substances.

A mixture does not have a fixed composition. For example, air is a mixture of gases. When students sit in a classroom, they use up oxygen and breathe out carbon dioxide and so the composition of the air in the room changes. We still call it 'air', but because air is a mixture its composition can change.



**3 a** Describe what a mixture of carbon, hydrogen and oxygen might look like.



**b** Describe how you would separate marbles from sand.



**4** Oxygen can be removed from air by cooling. Explain why this would not be possible if air were not a mixture.



## Melting points

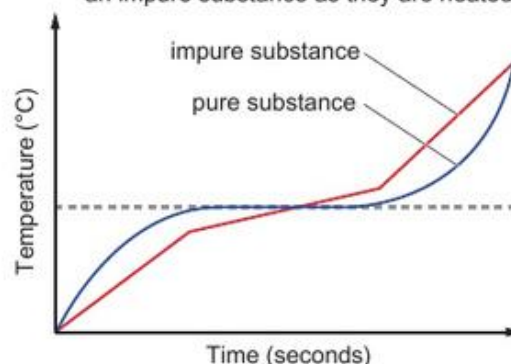
When a solid melts, its particles gain enough energy to overcome the weak forces of attraction between them. They move further away from one another and the solid becomes a liquid. The temperature at which this happens is the **melting point**. This is an example of a **physical property** (how a substance responds to forces and energy).

A pure substance has the same composition in every part of it, and so its physical properties are the same in every part. So, all of a pure substance will melt at the same temperature until all the substance has changed state. The melting point of pure gold is  $1063^{\circ}\text{C}$  and the melting point of oxygen is  $-218^{\circ}\text{C}$ .

- 5 What is the freezing point of pure oxygen?



How temperature changes in a pure substance and an impure substance as they are heated



C This sweet is a mixture and so does not have a sharp melting point.

D heating curves for a pure substance and a mixture

The sweet shown in photo C has a liquid centre. The whole sweet melts over a *range* of temperatures and not all the parts melt and become liquid at the same time. This is what happens in mixtures – they do not have fixed, sharp melting points.

Substance	Melting temperatures ( $^{\circ}\text{C}$ )
lead-tin alloy	183 to 258
argon	$-189$
carbon monoxide	$-205$

6 The table shows some melting temperatures.



a Identify which substances are mixtures and which are pure.



b Sketch a cooling curve for each of the three examples and explain their shapes.

### Exam-style question

Explain why mixtures melt over a range of temperatures but pure substances have precise melting points.

(2 marks)

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

S1 List the ways in which pure substances are different from mixtures.

### Extend

E1 A piece of gold jewellery is 750 on the fineness scale. Would you expect the jewellery to have a sharp melting temperature? Explain your answer.

# SC2b Filtration and crystallisation

Specification reference: C0.6; C2.7

## Progression questions

- How can filtration be used to separate mixtures?
- How can crystallisation be used to separate mixtures?
- What are the hazards and risks when separating mixtures by filtration and crystallisation?



**A** Some whales filter sea water with bristles (called baleen plates) to separate krill from the water.

- 1**
- a** Give one example of a mixture that can be separated by filtration.
- b** Explain how this mixture is separated by filtration.
- c** Describe another type of mixture that can be separated by filtration.

## Did you know?

Nearly 4 million tonnes of salt are solution mined in the UK each year.



**C** Salt can be produced by the evaporation of sea water.

Filters can be used to separate some mixtures. They let smaller pieces or liquids through but trap bigger pieces or **insoluble** substances.

Examples of **filtration** are to be seen all around us. Cars, vacuum cleaners and air-conditioning systems all have filters. Some whales use filters to feed. They open their mouths and take in water. When they close their mouths, they push out the water through filters. Small animals (such as krill) get stuck in the filters and are swallowed.

## Crystallisation

A **solution** is a mixture made of **solutes** (dissolved substances) in a liquid called the **solvent**. Solutes can be separated from a solution by evaporating the solvent to leave the solutes behind. This is called **crystallisation**. The process forms solid crystals of various sizes. If the crystals form slowly, the particles have longer to form an ordered pattern and will make larger crystals.



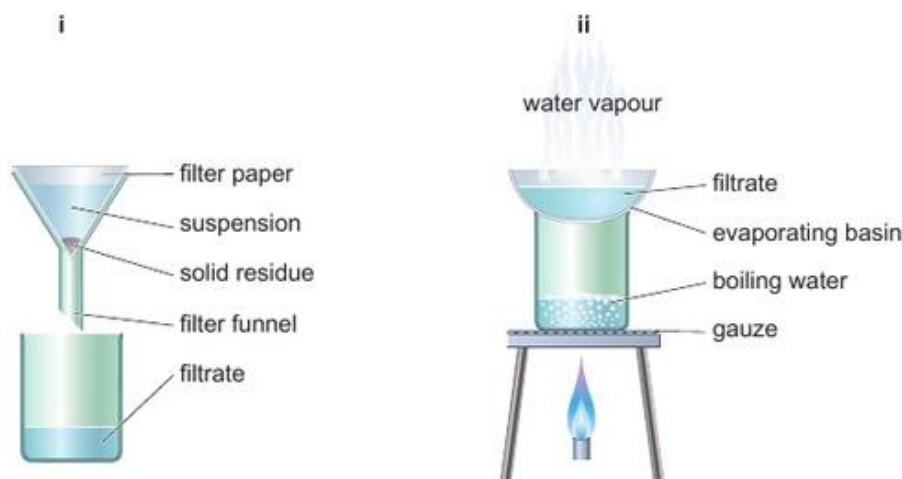
**B** Crystals in the Giant Crystal Cave in Mexico took over 500 000 years to form.

Table salt is produced from sea water, or is dug out of the ground or extracted using 'solution mining'. In this process water is pumped into layers of salt underground. The resulting salt solution is then heated, which evaporates the solvent and makes the solution more and more salty. Eventually it reaches a point where there is as much salt in the water as can possibly dissolve. This is a **saturated solution** and it contains the maximum amount of solute that can dissolve in that amount of solvent at that temperature. If more water evaporates and/or the solution cools, then some solute leaves the solution and salt crystals form.



## Filtration and crystallisation in the lab

To filter a solution in the laboratory, a filter funnel is lined with filter paper that has fine holes in it. The solvent and solute(s) pass through the fine holes to form the **filtrate**. Bits of insoluble substances cannot fit through the holes and so leave a **residue** in the filter paper. A Bunsen burner is then used to evaporate the filtrate carefully. Care must be taken not to overheat the solution once it is saturated, because hot crystals may spit out. Further heating may also cause crystals to change chemically.



D Laboratory apparatus for (i) filtration and (ii) crystallisation.

In a **risk assessment**, the **hazards** of doing an experiment are identified. A hazard is something that could cause harm. Then ways of reducing the **risk** (chance) of a hazard causing harm are considered.

During crystallisation, the risks from spitting can be reduced by wearing eye protection, removing the Bunsen burner before the solution is completely dry and/or using steam to heat the evaporating basin gently (as above).

- 7 When a mixture of rock pieces, salt and water is filtered, what will be found as the:



a filtrate



b residue?



- 8 a List two of the hazards when carrying out filtration and crystallisation.



b Explain how the risks from each of your hazards can be reduced.



- 2 Give the names of two mixtures that can be separated by crystallisation.

- 3 In the solution mining of salt, give the names of the:



a solvent



b solution



c solute.



- 4 When is a solution said to be 'saturated'?



- 5 Explain why crystals form during crystallisation.



- 6 Explain why the crystals in photo B are so big.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1 Explain how you would separate sand *and* salt from a mixture of the two.

### Extend

- E1 Scientists looking for new substances in plants grind up the plants with methanol. This solvent dissolves many plant compounds. However, methanol is flammable and toxic (especially if the vapour is inhaled). Large crystals can be made to help scientists work out what the compounds are made of. Explain how you would make plant-compound crystals using methanol.

### Exam-style question

Explain the difference between a risk and a hazard.

(2 marks)

# SC2c Paper chromatography

Specification reference: C2.7; C2.9; C2.10

## Progression questions

- How can chromatography be used to separate mixtures?
- What are the differences between mixtures and pure substances on a chromatogram?
- How do you calculate an  $R_f$  value?



**A** Experts restoring an old painting – they need to know what substances were mixed together to produce the paints used by the original artist.

Inks, paints and foods often contain mixtures of coloured compounds.

**Chromatography** can be used to find out which coloured compounds the mixture contains. The type of chromatography used to analyse the substances in old oil paintings requires expensive machinery.

**Paper chromatography** is a simpler technique that works because some compounds dissolve better in a solvent than others. When a solvent moves along a strip of paper, it carries the different substances in the mixture at different speeds, so they are separated. The solvent is called the **mobile phase**. The paper contains the **stationary phase**, through which the solvent and dissolved substances move. The paper with the separated components on it is called a **chromatogram**.

**5th** 1 **a** How many different compounds are in substance X in diagram B?

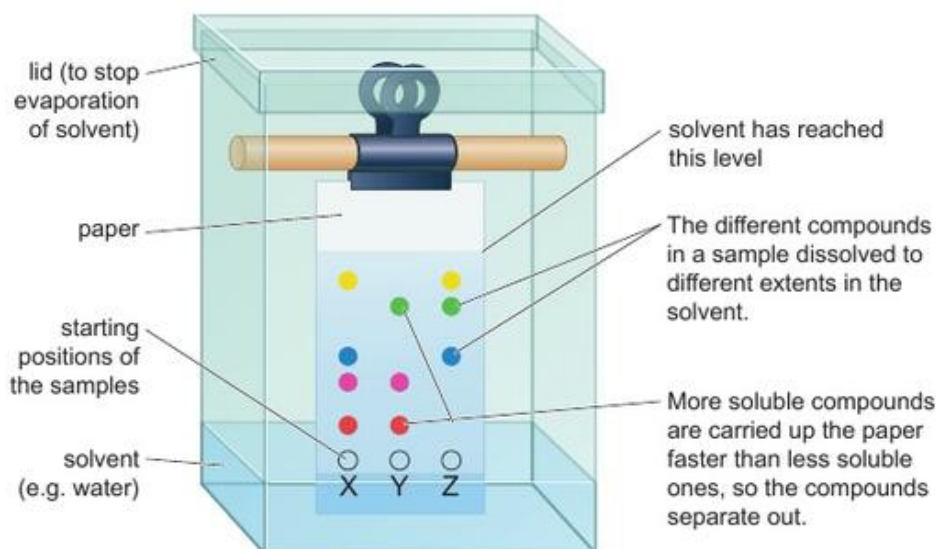
**6th** 1 **b** For mixture Y, explain why the green spot is higher than the red spot.

2 Look at diagram B again. Explain why:

**5th** 1 **a** the labels for substances X, Y and Z are written in pencil, not ink

**5th** 1 **b** the starting positions for the different substances are above the level of solvent in the container.

**6th** 3 One of the coloured compounds in diagram B has an  $R_f$  value of 0.1. Explain which compound this is likely to be.



### **B** paper chromatography

The  **$R_f$  value** is the distance the compound has risen divided by the distance the solvent has risen. Both measurements are made from the starting positions of the samples on the paper.

$$R_f = \frac{\text{distance moved by the spot}}{\text{distance moved by the solvent}}$$

The  $R_f$  value of a particular compound does not change if the chromatography conditions used remain the same.



**Worked example**

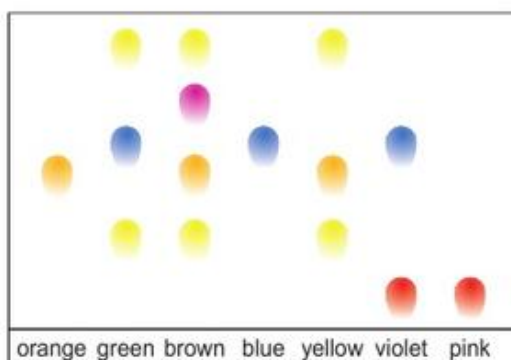
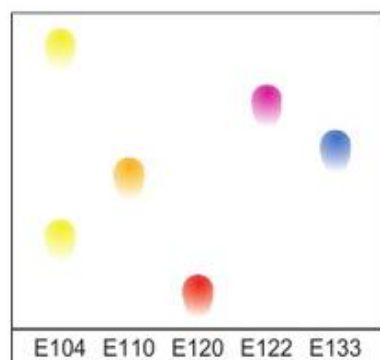
In diagram B, the pink spots have moved 4 cm and the solvent has moved 10 cm along the paper. Calculate the  $R_f$  value of this pink compound:

$$R_f = \frac{4}{10} = 0.4$$

A compound never rises as fast as the solvent, so  $R_f$  values are always less than 1. If you calculate an  $R_f$  value bigger than 1, you've made a mistake.

Paper chromatography can be used to:

- distinguish between pure and impure substances
- identify substances by comparing the pattern on the chromatogram with the patterns formed by known substances
- identify substances by calculating their  $R_f$  values.



**C** The chromatogram on the left was done using known substances. The chromatogram on the right shows that the orange and blue sweets contain single dyes.

**Did you know?**

In 1983, many national newspapers paid a lot of money to publish diaries allegedly written by Adolf Hitler. However, scientists used chromatography to analyse the inks in the diaries and found that they were not available during Hitler's lifetime – the diaries were fake.



**D** Chromatography can be used to help identify substances at crime scenes.

**Exam-style question**

Two dyes have the same  $R_f$  values when tested using chromatography. Explain whether this means they are the same dye or not. (3 marks)



**4** In diagram B, the yellow spots have moved 9 cm and the solvent has moved 10 cm. Calculate the  $R_f$  value of the yellow substance.

**5** In diagram C, the chromatogram on the left shows some food dyes found in sweets. The chromatogram on the right shows the results for some sweets.



**a** Which sweets contain just one dye?



**b** Which dyes are in the yellow sweets?



**c** What is the colour of the most soluble dye?

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

**S1** The police have taken four orange lipsticks from suspects. Explain the steps needed to find out if one of the lipsticks could have made a mark at a crime scene.

**Extend**

**E1** A laboratory produces a list of  $R_f$  values for food colourings. Explain why  $R_f$  values are used and what other information is needed for these  $R_f$  values to be useful.

# SC2d Distillation

Specification reference: C0.6; C2.7

## Progression questions

- What is distillation?
- How do simple distillation and fractional distillation differ?
- How would you reduce risks when carrying out a distillation experiment?



A a steam iron

Tap water contains dissolved minerals, especially in hard water areas – tap water is a **mixture**. For some jobs it is best to use pure water (such as for chemical analysis, in car-cooling systems and older steam irons). To make water pure we need to separate it from the dissolved solids. This is done by **distillation**.



**1 a** What happens to the water in a steam iron when you turn the iron on?



**b** Explain why some irons may not work well if you use ordinary tap water.



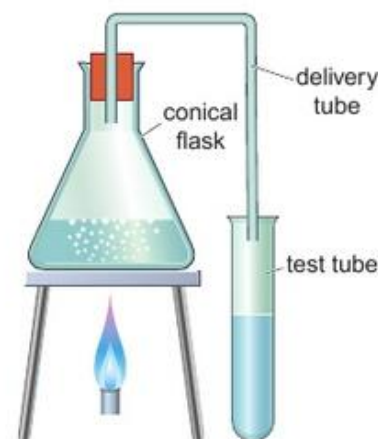
**2** In diagram B, what is the hazard if steam escapes from the tube?



**3** Suggest a way of improving a simple still so that more of the steam condenses back to water.

When mineral water **evaporates**, only the water turns to a gas (vapour). The solid minerals, which have much higher boiling points, are left behind. The water vapour (steam) is pure. If the vapour is then **condensed**, it turns back to liquid water again – the liquid water will now be pure. This combination of evaporation followed by condensation is called distillation. The apparatus used is called a **still**.

Diagram B shows a simple method for distilling. Water is heated in the conical flask and the vapour travels along the delivery tube, where it condenses. This method is not very efficient because much of the vapour is lost.



B a simple still

The type of still shown in diagram C is more efficient. The condenser keeps the tube cool, so that almost all of the vapour condenses and turns into a liquid.



**4** Explain how the still in diagram C can be used to purify water.

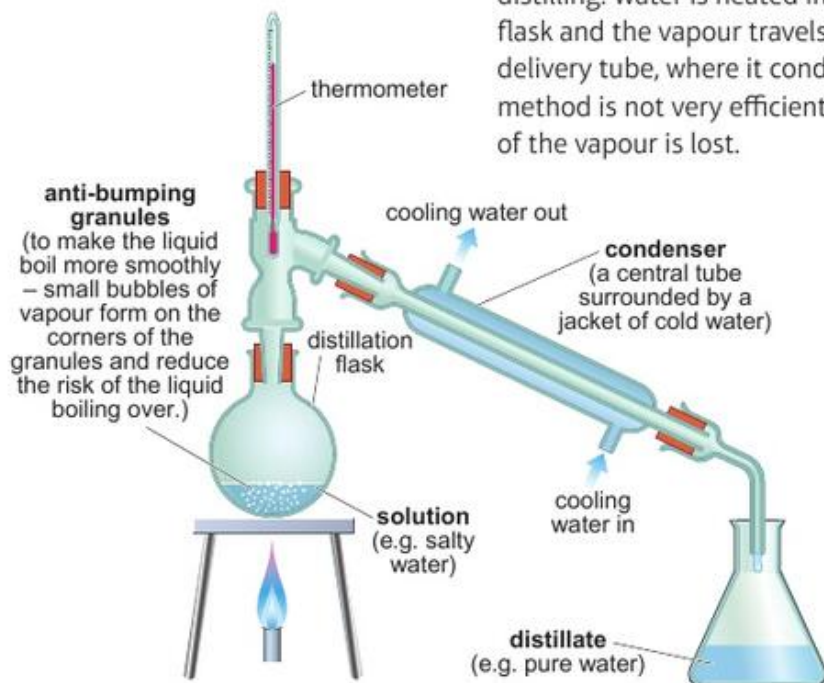
**5** Explain how:



**a** the condenser reduces the risk of hot vapour escaping



**b** the safety of the method is improved by using anti-bumping granules.



C distillation apparatus



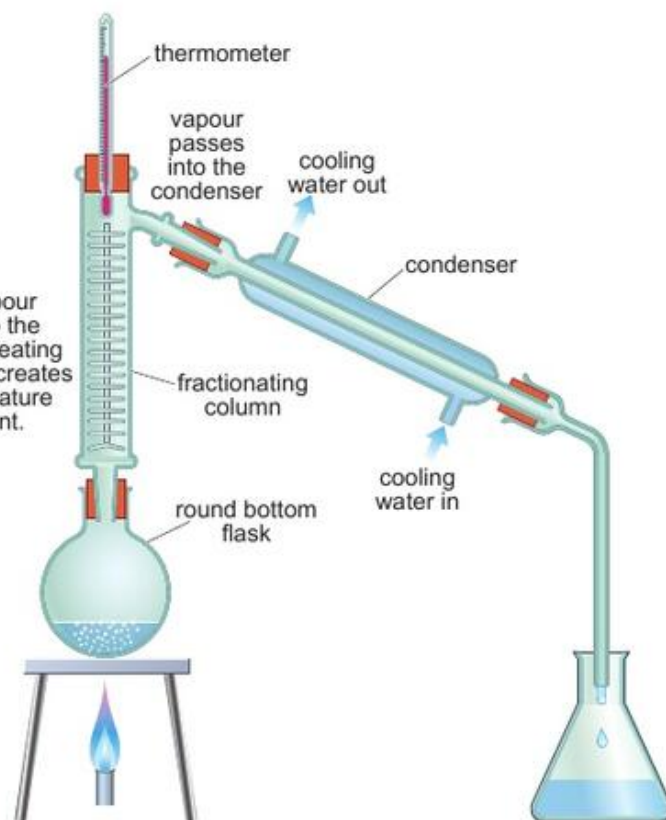
## Fractional distillation

Distillation can also be used to separate two or more liquids. This works because some liquids boil more easily than others. Liquids with lower boiling points evaporate more easily than others, and will turn into a vapour first. This is called **fractional distillation**, because the original mixture will be split into several parts, or fractions. The first fraction to be collected contains the liquid with the lowest boiling point. The fractions could be pure liquids, or may still be mixtures.

Fractional distillation can be used:

- to separate the different products in crude oil
- to make alcoholic drinks such as whisky and vodka
- to separate out the gases in the air, after the air has been cooled and turned into a liquid at  $-200^{\circ}\text{C}$ .

Diagram D shows how to separate liquids more efficiently. A column is fixed above the distillation flask. The hot vapour rises up the column. At first, the vapour condenses when it hits the cool glass and drips back down into the flask. As the column gradually heats up, there will be a temperature gradient – it will be hottest at the bottom and the temperature will drop as you go further up the column. The fraction with the lowest boiling point will reach the top of the column first and the vapour will then pass into the condenser. If you keep heating, fractions with higher boiling points will then rise up the column and can be collected later.



D distillation apparatus with a fractionating column

- 8<sup>th</sup>** 6 Compare and contrast simple and fractional distillation.
- 9<sup>th</sup>** 7 Explain why a liquid with a lower boiling point will reach the top of a fractionating column more quickly than one with a higher boiling point.

### Did you know?

The vacuum flask that we now use to keep drinks hot was originally used to keep liquid air *cold*. It was designed by James Dewar in 1892.

### Exam-style question

A student is asked to separate two liquids. Liquid A boils at  $100^{\circ}\text{C}$  and liquid B boils at  $65^{\circ}\text{C}$ . The student sets up a fractional distillation experiment, and after a few minutes a clear liquid is collected from the condenser. Explain which of the two liquids will be collected first. (2 marks)

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Explain what distillation is and how the distillation apparatus (the still) works. Use a labelled diagram to make your explanation clear.
- S2** Explain the safety precautions you need to take when carrying out distillation in a laboratory.

### Extend

- E1** Pure ethanol ('alcohol') boils at  $78.5^{\circ}\text{C}$ . Explain how a 50:50 mixture of ethanol and water can be separated by fractional distillation.
- E2** Suggest why the boiling point of the starting liquid will change with time.

# SC2d Core practical – Investigating inks

Specification reference: C2.11

## Aim

Investigate the composition of inks using simple distillation and paper chromatography.



**A** Fountain pen ink is available as a washable ink.

Ink is a mixture of coloured substances dissolved in a liquid solvent. Ink that appears to be a single colour, such as black, may contain two or more substances with different colours. Permanent inks do not run if the paper becomes wet, a useful property if you drop your homework in a puddle. Washable inks separate into their different colours if the paper gets wet, but may be removed if spilt on clothing.

## Your task

You are going to use simple distillation to separate a sample of the solvent in some ink. You will also use paper chromatography to separate the coloured substances in samples of ink.

## Method

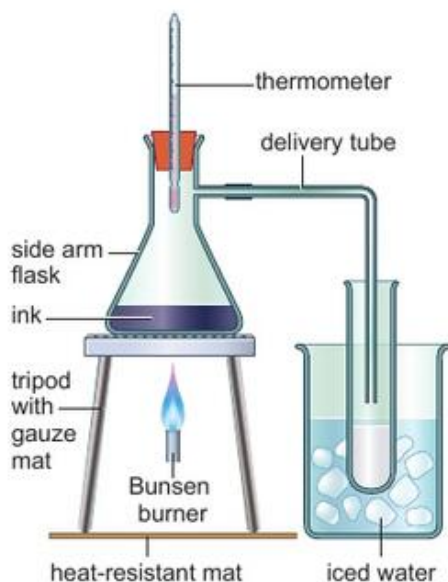
### Simple distillation

Wear eye protection.

- A** Set up your apparatus so that the ink is in a flask, and its vapours can be led away to be condensed. Diagram B shows some typical apparatus but yours may be different.
- B** Heat the flask of ink using a Bunsen burner, making sure the ink simmers gently and does not boil over into the delivery tube.
- C** Continue heating until you have collected a few cm<sup>3</sup> of distillate (distilled solvent).
- D** Note the maximum temperature obtained.

### Paper chromatography

- E** Draw a pencil line on a piece of chromatography paper, about 2 cm from the bottom.
- F** Add a small spot of ink to the pencil line.
- G** Add water to a container to a depth of about 1 cm.
- H** Place the paper into the container. Make sure the paper is supported so that it does not slump into the water when it becomes damp. Allow the water to travel through the paper.
- I** Take the paper out before the water reaches the top. Immediately mark the position of the solvent front using a pencil, then leave the paper to dry.
- J** Measure the distance travelled by the water from the pencil line, and the distances travelled by each coloured substance.
- K** Calculate the  $R_f$  value for each coloured substance.

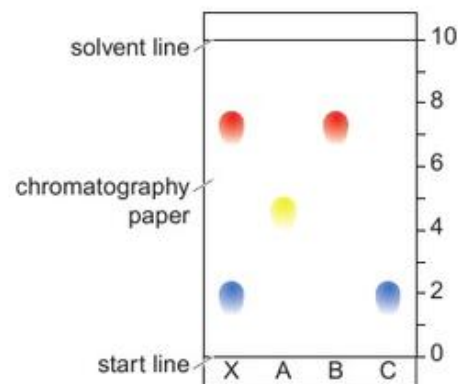


**B** separating ink using simple distillation



## Exam-style questions

- 1 Explain the function of the beaker of iced water in diagram B.  
(2 marks)
- 2 A student carries out simple distillation on a sample of blue ink.
  - a Predict how the appearance of the ink changes, and give a reason for your answer.  
(2 marks)
  - b During the experiment, hot liquid solvent drips from the bulb of the thermometer. Suggest an explanation for a temperature rise from 83 °C to 100 °C as this happens.  
(1 mark)
- 3 Explain why simple distillation allows a pure solvent to be separated from a solution.  
(3 marks)
- 4 A student distils a sample of ink. Devise a simple method to show that the liquid collected is pure water. Include the expected results in your answer.  
(3 marks)
- 5 Propanone is a flammable solvent. A student carries out paper chromatography of ink using propanone.
  - a Identify the mobile phase and the stationary phase in her experiment.  
(2 marks)
  - b Explain one precaution necessary to control the risk of harm in her experiment.  
(1 mark)
  - c Suggest an explanation for why the level of the propanone should be below the ink spot on the paper at the start.  
(1 mark)
- 6 Explain how paper chromatography separates coloured substances in ink.  
(3 marks)
- 7 A student uses paper chromatography to analyse four samples of ink (X, A, B and C). Diagram C shows his results.
  - a Describe what the results tell you about ink sample X.  
(2 marks)
  - b Calculate the  $R_f$  value of the substance in ink B.  
(2 marks)
- 8 A student uses paper chromatography to analyse the dyes present in a sample of ink. She adds a sample of the ink and four dyes (1, 2, 3 and 4) to the paper. Table D shows her results for the dyes.
  - a Explain which dye is the most soluble in the solvent used by the student.  
(2 marks)
  - b Explain whether each dye is a pure substance.  
(2 marks)
  - c Suggest an explanation for why a mixture of dyes 1 and 4 may appear as a single green spot in a paper chromatogram.  
(2 marks)



C

Dye	Spot colour	$R_f$
1	yellow	0.10
2	red	0.35
3	green	0.67
4	blue	0.12

D

# SC2e Drinking water

Specification reference: C2.8; C2.12

## Progression questions

- How would you choose which method to use to separate a mixture?
- How is drinking water produced?
- Why must water used in chemical analysis be pure?



**A** Personal water purifiers filter water to make it safer to drink.



**1** Explain how pure water is produced using the apparatus shown in diagram B.



**2** Suggest why the simple distillation of sea water may be used to provide drinking water in oil-rich coastal countries.

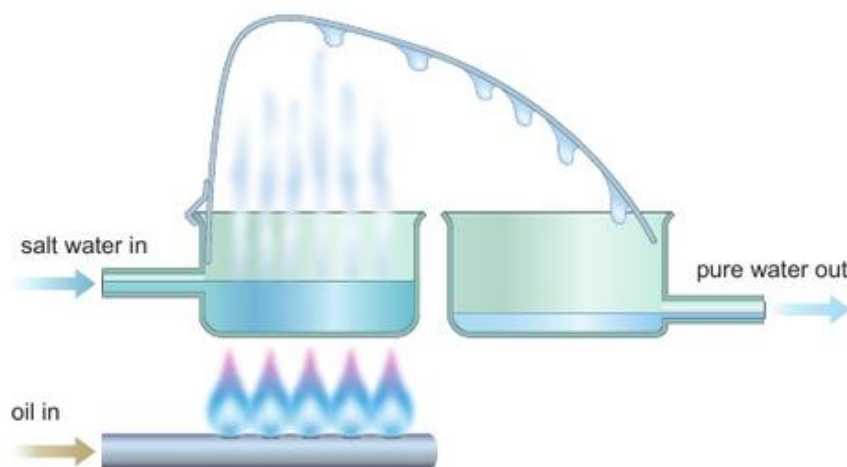


**C** A cloudy white precipitate forming during a chemical analysis.

About 97% of the Earth's water is in the oceans. The concentration of dissolved salts in sea water is far too high for us to drink safely. Producing pure water from sea water is called **desalination** and can be achieved using **simple distillation**.

## Purifying sea water

Water is separated from dissolved salts using simple distillation. Sea water is heated so that water vapour leaves it quickly. This vapour is then cooled and condensed, forming water without the dissolved salts.



**B** Simple distillation of sea water using oil as a fuel.

A lot of energy must be transferred to sea water during simple distillation, so it is not usually a suitable method for producing large volumes of drinking water. It is mainly carried out on a large scale where energy resources are cheap or plentiful, and where there is an abundant supply of sea water.

## Water for chemical analysis

**Chemical analysis** involves using chemical reactions or sensitive machines to identify and measure the substances in a sample. The water used for chemical analysis should not contain any dissolved salts, otherwise incorrect results will be obtained. Tap water contains small amounts of dissolved salts, which may react to form unexpected cloudy **precipitates**. These may hide the correct result of the analysis. Also, the machines used for analysis may detect the salts, again leading to an incorrect conclusion.



**3** Explain why distilled water is more suitable than tap water for doing a chemical analysis.

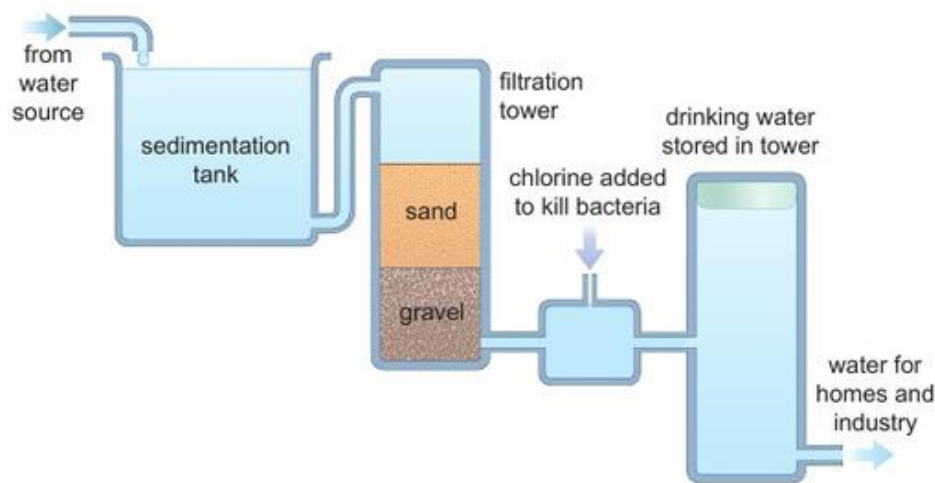


## Water for drinking

In the UK, the raw material for producing drinking water comes from rivers, lakes or **aquifers** (underground rocks containing groundwater). The water in these sources is often stored in reservoirs, which are artificial lakes produced by building a dam across a valley. Fresh water from these sources contains:

- objects such as leaves and twigs
- small insoluble particles such as grit and silt
- soluble substances, including salts, pesticides and fertilisers
- bacteria and other microorganisms that may be harmful to health.

Different steps are needed to deal with these impurities. They include screening using a sieve, **sedimentation** (in which small particles are allowed to settle out) and filtration using tanks containing beds of sand and gravel. Chlorine is added in a process called **chlorination**. Chlorine kills microorganisms in the treated water.



**D** These are the main stages in treating fresh water to make it safe to drink.

- 4** **5** a Describe how water is treated to deal with leaves and twigs, grit and silt, and with microorganisms.
- 5** **b** Identify the stage missing from diagram D and draw a labelled diagram to show it.
- 6** Suggest why chemical reactions, rather than separation methods, are used to remove harmful substances dissolved in drinking water in the UK.

### Exam-style question

Fresh water is treated to make it safe to drink. Soluble and insoluble substances are removed during this treatment, and chlorine is added to kill harmful microorganisms. State two reasons why samples of the treated water are tested regularly. (2 marks)



- 4** Explain why it may not be safe to drink water straight from a river.

### Did you know?

Only about 2.5% of the Earth's water is fresh water. Of that, only 0.3% is in rivers and lakes – the rest is in icecaps, glaciers and ground water.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Draw flowcharts to describe two ways in which water can be made fit to drink.

### Extend

- E1** A bottle of water has a label saying 'Suitable for chemical analysis'. Describe how this water has been produced.
- E2** Explain how you would check to see if this water really is suitable for analysis.

# Methods of separating and purifying substances

A runny green mixture contains three compounds (**X**, **Y** and **Z**).

Compound	Melting point (°C)	Boiling point (°C)	State at room temperature	Notes
X	2435	4000	solid	not soluble in <b>Y</b> or <b>Z</b>
Y	-126	97.4	liquid	soluble in <b>Z</b>
Z	-114	78.4	liquid	soluble in <b>Y</b>

Plan a method to separate out **X**, **Y** and **Z**. Use the information in the table above and explain why you have suggested each step. (6 marks)

## Student answer

Compound X is solid, and it is insoluble in the two liquids (Y and Z). This means it can be separated from them by filtration [1]. When X is collected in the filter paper it will be wet with the two liquids, so it should be dried in a warm oven. The boiling points of the two liquids are well below the melting point of compound X, so compound X will not melt in the oven [2].

The filtrate will be a mixture of compounds Y and Z. These can be separated using fractional distillation because their boiling points are different [3]. When the mixture is heated, compound Z will distil off first because it has the lower boiling point [4], leaving compound Y behind.

[1] This explains why compound X can be separated by filtration.

[2] The answer uses information from the table to explain why drying compound X will work.

[3] This makes it clear which physical property fractional distillation depends on.

[4] It is clear which liquid will be collected first, and why.

## Verdict

This is a strong answer. It shows good knowledge and understanding of separation methods and uses correct scientific language.

The answer is organised logically, in the order that the practical would be carried out. Each step of the practical is linked to a scientific explanation.

### Exam tip

If you are given a table of data or a graph in a question, make sure you use information from it in your answer.





## Paper 1 and Paper 2

### SC3 Atomic Structure

This image courtesy of Prof Richard Palmer, University of Birmingham, shows palladium atoms on a base of carbon. According to the scientists who produced it, Zhiwei Wang and David Pearmain, although they had watched with love, they had nothing to do with the spontaneous formation of the heart shape. Unfortunately this atomic valentine, being only 8 nanometres (0.000000008 metres) across, is far too small to see even with the strongest light microscope. It may, however, help to explain the nature of matter, which is central to understanding the properties of materials and the chemical reactions that form new substances. In this unit you will find out more about atoms and their structure.

#### The learning journey

Previously you will have learnt at KS3:

- about the particle model of matter
- how Dalton's ideas about atoms helped to explain the properties of matter
- how elements are arranged in the periodic table.

In this unit you will learn:

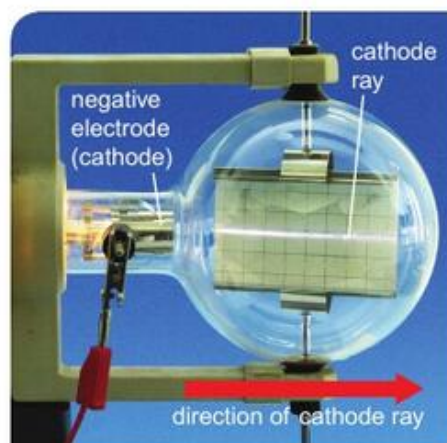
- how our ideas about atoms have changed
- what a relative atomic mass is
- **H** how to calculate relative atomic mass for an element.

# SC3a Structure of an atom

Specification reference: C1.1; C1.2; C1.3; C1.4; C1.5

## Progression questions

- How has the model of the atom changed over the last 200 years?
- How do the parts of atoms compare with each other?
- Why do atoms have no overall charge?



A a cathode ray tube

In 1805 the English chemist John Dalton (1766–1844) published his atomic theory that said:

- all matter is made up of tiny particles called **atoms**
- atoms are tiny, hard spheres that cannot be broken down into smaller parts
- atoms cannot be created or destroyed
- the atoms in an **element** are all identical (but each element has its own type of atom).

Dalton's ideas helped to explain some of the properties of matter. However, experiments towards the end of the nineteenth century suggested that atoms contain even smaller particles.

When a high voltage is applied to a glass tube that has most of the air removed, glowing rays are seen. Some scientists thought that these 'cathode rays' were atoms leaving the negative electrode. In 1897, JJ Thomson (1856–1940) investigated the mass of the particles in the rays and found that they were about 1800 times lighter than the lightest atom (hydrogen). Cathode rays, therefore, did not contain atoms but **subatomic particles**, which we now call **electrons**.

## The structure of atoms

Scientists have now worked out that atoms are made up of electrons together with heavier subatomic particles called **protons** and **neutrons**. All these particles have very, very small masses and electric charges. So, rather than use their actual masses and charges, it is easier to describe them by looking at their **relative masses** and **relative charges** compared to a proton. For example, if we say the mass of a proton is '1' then anything else that has the same mass is also '1'.

### Did you know?

The actual mass of a proton is 0.00000000000000000000000167 g ( $1.67 \times 10^{-24}$  g).

Subatomic particle	Relative charge	Relative mass
proton	+1 (positive)	1
electron	-1 (negative)	1/1835 (negligible)
neutron	0 (no charge)	1

### B relative masses and relative charges of subatomic particles

At the centre of all atoms is a tiny **nucleus** containing protons and neutrons. This is surrounded by fast moving electrons arranged in **electron shells** at different distances from the nucleus.

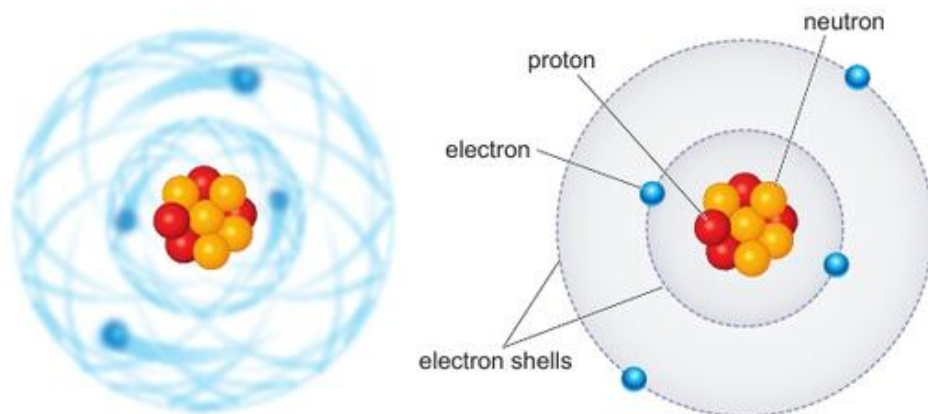
- 7<sup>th</sup> 1 What are atoms?
- 8<sup>th</sup> 2 Which of Dalton's ideas about particles is supported by the image of palladium atoms on the previous page?

- 7<sup>th</sup> 3 Which subatomic particle has the lowest mass?



Atoms in elements always have equal numbers of protons and electrons and so have no overall charge, because the charges cancel out.

Diagram C shows two ways of modelling a beryllium atom. The three-dimensional model attempts to show how we imagine electrons to move.



**C** The 'target diagram' on the right shows the arrangement of the electrons more clearly.

Models of atoms help us to understand their structure – but most models don't really give a correct impression of scale. The overall diameter of an atom can be 100 000 times the diameter of its nucleus.



**D** If an atom could be made the size of the Lord's cricket ground, its nucleus would be about the size of this dot ●. Most of an atom is empty space.

### Exam-style question

Compare the modern model of an atom to the atomic model proposed by John Dalton in 1805. (2 marks)

- 4** How many protons, neutrons and electrons are in a beryllium atom?
- 5** A lithium atom has 3 protons, 4 neutrons and 3 electrons.
- a** Draw a diagram of this atom.
- b** Why is this atom neutral?
- c** How many electrons would be in an atom that has 17 protons?

### Checkpoint

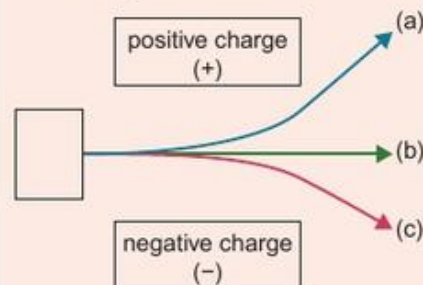
How confidently can you answer the Progression questions?

### Strengthen

**S1** Draw an atom and label it to describe the arrangement and properties of its subatomic particles.

### Extend

**E1** Figure E (below) shows what happens when the three subatomic particles are fired through an electric field.



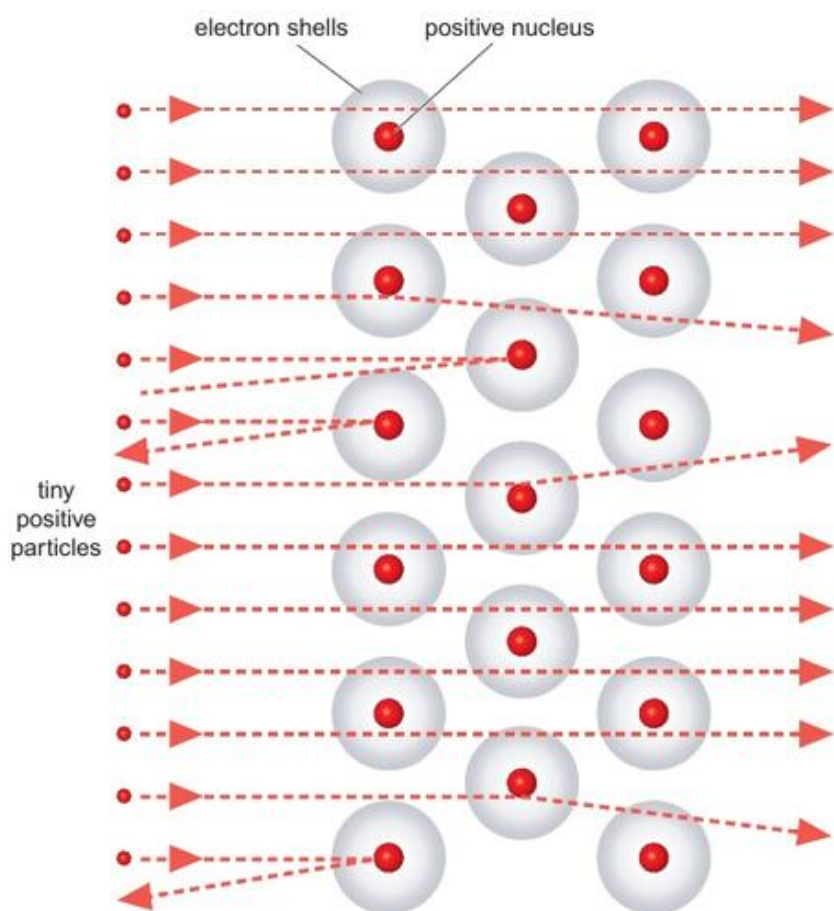
Name each particle a, b and c. Explain your answer.

# SC3b Atomic number and mass number

Specification reference: C1.6; C1.7; C1.8; C1.10

## Progression questions

- Why is most of the mass of an atom found in its nucleus?
- What does the atomic number tell you about an element?
- How can you calculate the numbers of protons, neutrons and electrons in atoms?



A Rutherford's scattering experiments suggested a nuclear atomic model.

## The nuclear atom

In 1909, Ernest Rutherford (1871–1937) was working with others to investigate the structure of atoms. In one experiment, tiny positive particles were fired at a thin gold foil. To everyone's surprise most of the particles passed straight through the gold foil, with a few being deflected and a very small number bouncing back. Rutherford explained this by suggesting that atoms are mostly empty space, with a small positive central nucleus that contains most of the mass.



1 a Where is most of the mass of an atom?



b Explain how the experiment in diagram A suggests that atoms are mostly empty space.

## Atomic number

The elements in the **periodic table** were originally placed in order of the masses of their atoms. However, this caused some elements to be grouped with others that had very different properties. So a few elements

were swapped round to make sure that those with similar properties were grouped together, even if it meant that they were no longer in the correct order of mass.

Experiments by Henry Moseley (1887–1915) in 1913 confirmed that the rearranged order of elements in the table was actually correct. He showed that they were in order of the amount of positive charge in the nucleus. The proton was discovered about five years later. The modern periodic table places the elements in order of the number of protons in their atoms. This is the **atomic number** and it is this that defines an element – all the atoms of a particular element have the same unique atomic number.

## Did you know?

Henry Moseley was killed during the First World War. As a result of Moseley's death, other important scientists were restricted from serving in front-line roles.



2 Carbon has an atomic number of 6. How many protons does it have?



3 Use a periodic table to find:



a the number of protons and electrons in atoms of:

- i nitrogen                      ii potassium



b two elements whose atomic mass order does not match their atomic number order.

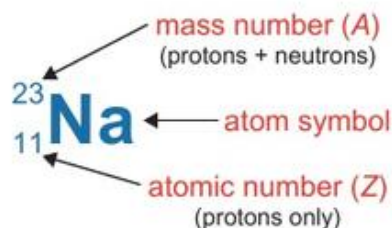


4 In terms of structure, what do all atoms of a certain element have in common?

## Mass number

The mass of an electron is described as 'negligible' – it is so small that it can be ignored. This explains why the nucleus of any atom contains nearly all its mass. For this reason the total number of protons and neutrons in an atom is called its **mass number**.

A mass number is represented by the symbol  $A$  and an atomic number by the symbol  $Z$ . These numbers are written next to an element's symbol as shown in diagram B.



**B** This is how scientists write the atomic number and the mass number for a sodium atom. It shows that the atom contains 11 protons and 12 neutrons in its nucleus.



5 How many protons, neutrons and electrons are in the atom  $^{27}_{13}\text{Al}$ ? Explain how you worked out your answer.



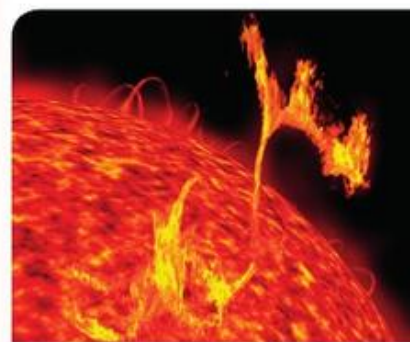
6 A manganese atom has 25 protons, 30 neutrons and 25 electrons. Show this information using the form shown in diagram B.



7 Look at the 'Did you know?' box on this page. Which subatomic particle does a hydrogen atom *not* have? Explain your reasoning.

## Did you know?

Just two elements make up most of our Universe – about 74% is hydrogen  $^1_1\text{H}$  and 24% is helium  $^4_2\text{He}$ . These are also the two simplest atoms in the periodic table. The Sun releases energy by converting hydrogen atoms into helium atoms. Every second, over 600 million tonnes of hydrogen is converted to helium.



**C**

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** An atom can be represented in the form  $^{65}_{29}\text{Cu}$ . What does this tell you about this atom?

## Extend

**E1** Formulae can be written to connect the atomic number, mass number, and numbers of protons, electrons and neutrons in atoms.

For example:

*atomic number = protons*

Write formulae that connect the other four numbers.

## Exam-style question

Complete the table below

(2 marks)

Atom	atomic number	mass number	number of protons	number of neutrons	number of electrons
X	90	222	90	132	(i)
Y	88	(ii)	88	134	88

# SC3c Isotopes

Specification reference: C1.9; C1.10; C1.11; **H** C1.12

## Progression questions

- How can you describe and identify isotopes of elements?
- Why are the relative atomic masses for some elements not whole numbers?
- **H** How do you calculate the relative atomic mass of an element?

**7<sup>th</sup>** **1 a** Write the name of each of the three lithium isotopes in diagram A.

**8<sup>th</sup>** **b** Describe each isotope in the format  ${}^A_Z\text{X}$  (X is the element's symbol).

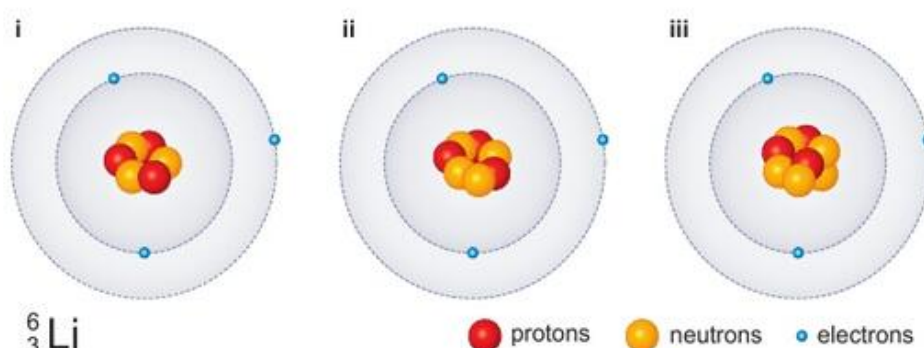
## Did you know?

In 1945, at the end of the Second World War, two 'atomic bombs' were dropped on the Japanese cities of Hiroshima and Nagasaki. The bombs used nuclear fission and killed at least 129 000 people.



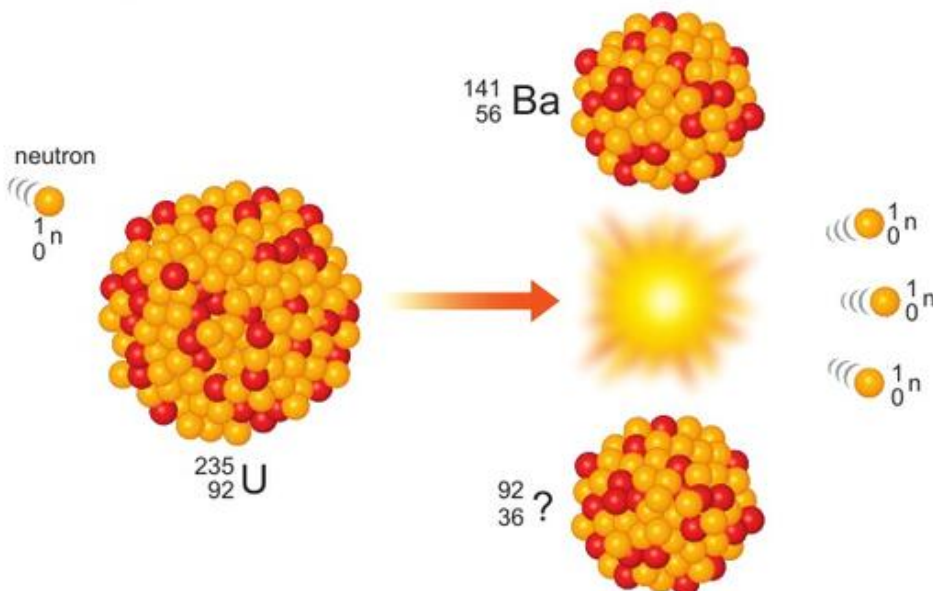
**B**

In 1932, James Chadwick (1891–1974) discovered the neutron. His discovery explains why some atoms of the same element have different masses. These atoms are known as **isotopes** – they have the same atomic number but different mass numbers. We refer to a specific isotope by adding its mass number to the element's name. The isotope on the left of diagram A is lithium-6.



**A** Isotopes of the same element are chemically identical because they have the same number of protons and electrons.

Understanding neutrons led to the discovery of nuclear energy. By firing neutrons at a uranium isotope,  ${}^{235}_{92}\text{U}$ , it was discovered that a nucleus can be split (**nuclear fission**). This produces new elements and transfers large amounts of energy. Nuclear power stations use the energy from nuclear fission to produce electricity.



**C** Nuclear fission can be started by firing neutrons at atoms of uranium-235.

**7<sup>th</sup>** **2 a** State the numbers of protons and neutrons in an atom of uranium-235.

**7<sup>th</sup>** **b** State the number of protons, neutrons and electrons in the barium isotope formed in diagram C.

**7<sup>th</sup>** **c** Identify the other product. Give your answer using the format  ${}^A_Z\text{X}$ .

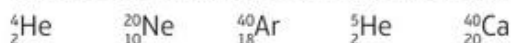


## Relative atomic masses

The mass of an atom is incredibly small, so we measure their masses *relative to* (compared to) an atom of carbon-12. This isotope is used as a standard and given a mass of exactly 12. The masses of all other atoms are compared to this. For example, the mass of helium-4 is one-third of that of carbon-12 and so its relative mass is 4.

**3** How many  ${}^4_2\text{He}$  atoms would be needed to balance the mass of the  ${}^{24}_{12}\text{Mg}$  atom in diagram D?

**4** Use the list of isotopes to answer the questions that follow.



Which two isotopes have the same:

**a** mass                      **b** chemical properties?

**5** Write the mass ratios, in their simplest form, for the isotopes:

**a**  ${}^{20}_{10}\text{Ne}$  and  ${}^{40}_{18}\text{Ar}$       **b**  ${}^4_2\text{He}$  and  ${}^{20}_{10}\text{Ne}$

The relative mass of an isotope is its mass number. For example, chlorine has two isotopes,  ${}^{35}_{17}\text{Cl}$  and  ${}^{37}_{17}\text{Cl}$ , and their relative masses are 35 and 37, respectively.

All elements exist as mixtures of isotopes. We use this idea to calculate an element's **relative atomic mass (RAM)** – the symbol is  $A_r$ . A relative atomic mass is the **mean** mass of an atom of an element compared with carbon-12. It takes into account all the isotopes of the element and the amounts of each. RAMs are not whole numbers (for example the  $A_r$  of chlorine is 35.5) but most values are commonly rounded to whole numbers. The RAM of an element and its atomic number are shown on the periodic table.

**H**

The abundances (overall proportions) of the two isotopes of chlorine are 75% of  ${}^{35}_{17}\text{Cl}$  and 25% of  ${}^{37}_{17}\text{Cl}$ . We calculate the relative atomic mass of chlorine as follows.

$$\begin{aligned} \text{If we take 100 atoms,} \\ \text{the relative atomic mass} &= \frac{\text{total mass of the atoms}}{\text{the number of atoms}} = \frac{(75 \times 35) + (25 \times 37)}{100} \\ &= \frac{2625 + 925}{100} = \frac{3550}{100} \\ A_r &= 35.5 \end{aligned}$$

### Exam-style question

Neon gas has a relative atomic mass of 20.2 and is made up of two atoms:  ${}^{20}_{10}\text{Ne}$  and  ${}^{22}_{10}\text{Ne}$ . Explain which of these isotopes is the most abundant.

(3 marks)



**D** An atom of the isotope  ${}^{24}_{12}\text{Mg}$  has twice the mass of  ${}^{12}_6\text{C}$  and so has a relative mass of 24.

**6** What does the relative atomic mass of an element tell you?

**7** **H** Copper has two isotopes – 69% is  ${}^{63}_{29}\text{Cu}$  and 31% is  ${}^{65}_{29}\text{Cu}$ . Calculate the RAM of copper. Give your answer to 1 decimal place.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

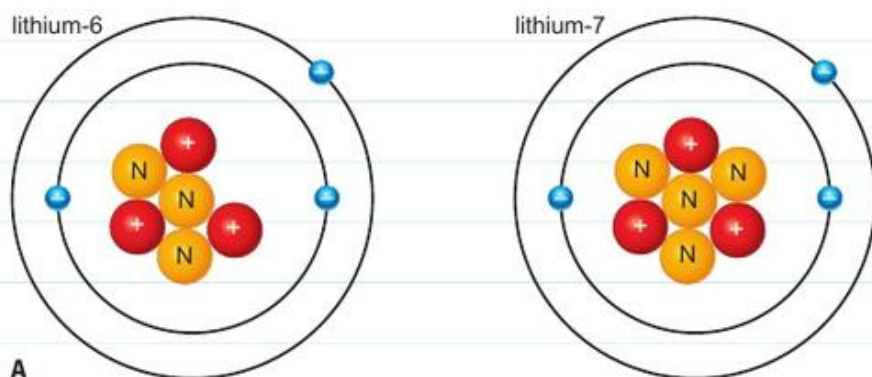
**S1** Describe, with examples, the similarities and differences between isotopes of the same element.

### Extend

**E1** **H** Describing each step, work out the relative atomic mass of magnesium – it has the composition 79%  ${}^{24}_{12}\text{Mg}$ , 10%  ${}^{25}_{12}\text{Mg}$  and 11%  ${}^{26}_{12}\text{Mg}$ .

## Isotopes

The diagram below shows the structure of two isotopes of lithium.



Using these lithium atoms as examples, explain the similarities and differences in the properties of different isotopes of the same element. (6 marks)

### Student answer

Atoms that are isotopes of the same element have the same atomic numbers but different mass numbers [1]. This means they have the same number of protons and electrons but different numbers of neutrons [2].

Atoms that are isotopes will chemically react in the same way [3] but be different in structure [4].

[1] The answer gives a basic definition of isotopes.

[2] This is a good description of the similarities and differences in the number of subatomic particles, but it doesn't use the lithium isotopes to illustrate the examples.

[3] This would be better if the answer explained *why* atoms of different isotopes react in the same way.

[4] This just repeats the earlier statement about differences, without adding any further detail.

### Verdict

This is an acceptable answer. It contains a basic definition of isotopes, and a description of their similarities and differences in terms of structure. The answer also notes that the isotopes will have the same chemical properties. There is some linking of scientific ideas and a basic logical structure.

This answer could be improved by referring to the examples given in the question. It would have been better if the answer had included clearer definitions of 'atomic number' and 'mass number'. The answer should also link together some more scientific ideas. For example, it could have linked reactivity with electron configuration (to explain why isotopes of the same element have the same reactivity).

### Exam tip

If an exam question asks you to use examples that are given, you must include them in your answer.



# Paper 1 and Paper 2

## SC4 The Periodic Table

There are over 100 known elements. The modern periodic table is a chart that arranges these elements in a way that is useful to chemists. Thanks to the periodic table, chemists can make sense of patterns and trends, which lets them predict the properties of elements. This works even if only a few atoms of an element exist. The periodic table shown here includes photos of most of the elements. If an element is very rare or difficult to obtain, it shows a photo of the relevant scientist or research laboratory involved in discovering or naming the element, or a diagram representing the arrangement of its electrons.

### The learning journey

Previously you will have learnt at KS3:

- about chemical symbols for elements
- that Dmitri Mendeleev designed an early periodic table
- about periods and groups in the periodic table
- about metals and non-metals, their properties and their positions in the periodic table.

In this unit you will learn:

- how Mendeleev arranged the elements known at the time in a periodic table
- how Mendeleev predicted the existence and properties of undiscovered elements
- how Henry Moseley helped to confirm Mendeleev's ideas
- how the elements are arranged in the modern periodic table
- how to use the periodic table to predict and model the arrangement of electrons in atoms.



# SC4a Elements and the periodic table

Specification reference: C0.1; C1.13; C1.14

## Progression questions

- What are the symbols of some common elements?
- How did Mendeleev arrange elements into a periodic table?
- How did Mendeleev use his table to predict the properties of undiscovered elements?



**A** Towels come in all sorts of sizes, colours and patterns, just as the elements have different properties. How would you organise them?

The Russian chemist Dmitri Mendeleev (1834–1907) faced a problem early in 1869. He was busy writing the second volume of his chemistry textbook and could not decide which elements it made sense to write about next. His solution was to construct a table that led to the **periodic table** we know today.

## Organising elements

Chemists had discovered 63 elements by 1869, and they were keen to organise them in a helpful way. Mendeleev arranged these elements in order of increasing **relative atomic masses** (called atomic weights then). Unlike other chemists who had tried this before, Mendeleev did not always keep to this order, and he left gaps in his table.

Mendeleev sometimes swapped the positions of elements if he thought that better suited their **chemical properties** and those of their compounds. For example, fluorine, chlorine, bromine and iodine are non-metals that do not easily react with oxygen, whereas tellurium is a metal that burns in air to form tellurium dioxide. Iodine has a lower relative atomic mass than tellurium, so Mendeleev should have placed it before tellurium according to this **physical property**. Instead, he placed iodine after tellurium so that it lined up with fluorine, chlorine and bromine (elements with similar chemical properties to iodine). Even though Mendeleev used the most accurate relative atomic masses then available, he justified this swap by stating that the value for tellurium must be incorrect.

Mendeleev assumed that elements would continue to be discovered, so he left gaps for them. This helped him to position the existing elements so that vertical columns contained elements with increasing relative atomic mass, and horizontal rows contained elements with similar chemical properties.

ОПЫТЪ СИСТЕМЫ ЭЛЕМЕНТОВЪ,  
ОСНОВАННОЙ НА ИХЪ АТОМНОМЪ ВѢСѢ И ХИМИЧЕСКОМЪ СХОДСТВѢ.

				Ti=50	Zr=90	?=180.
				V=51	Nb=94	Ta=182.
				Cr=52	Mo=96	W=186.
				Mn=55	Rh=104,4	Pt=197,1.
				Fe=56	Ru=104,4	Ir=198.
				Ni=Co=59	Pd=106,6	Os=199.
H=1				Cu=63,4	Ag=108	Hg=200.
	Be=9,4	Mg=24		Zn=65,2	Cd=112	
	B=11	Al=27,3		?=68	Ur=116	Au=197?
	C=12	Si=28		?=70	Sn=118	
	N=14	P=31		As=75	Sb=122	Bi=210?
	O=16	S=32		Se=79,4	Te=128?	
	F=19	Cl=35,5		Br=80	I=127	
Li=7	Na=23			K=39	Rb=85,4	Cs=133
				Ca=40	Sr=87,6	Ba=137
				?=45	Ce=92	Pb=207.
				?Er=56	La=94	
				?Yt=60	Di=95	
				?In=75,6	Th=118?	

**B** Mendeleev's published 1869 table. The question mark after the relative atomic mass of tellurium, Te, shows that he thought this was incorrect. He swapped the positions of iodine and tellurium so that iodine ended up in the same line as other elements with similar properties.



**1** What information about the elements did Mendeleev use to produce his first table?



**2** Explain why Mendeleev swapped the positions of iodine and tellurium in his table.

## Making predictions

Mendeleev continued to work on his table. By 1871, he had settled on a table in which elements with similar properties were arranged into vertical columns, just as today.



Series	Group 1	Group 2	Group 3	Group 4	Group 5	Group 6	Group 7	Group 8
1	H 1							
2	Li 7	Be 9.4	B 11	C 12	N 14	O 16	F 19	
3	Na 23	Mg 24	Al 27.3	Si 28	P 31	S 32	Cl 35.5	
4	K 39	Ca 40	? 44	Ti 48	V 51	Cr 52	Mn 55	Fe 56 Ni 59 Co 59 Cu 63
5	(Cu 63)	Zn 65	? 68	? 72	As 75	Se 78	Br 80	
6	Rb 85	Sr 87	Y 88	Zr 90	Nb 94	Mo 96	? 100	Ru 104 Rh 104 Pd 106 Ag 108
7	(Ag 108)	Cd 112	In 113	Sn 118	Sb 122	Te 125	I 127	
8	Cs 133	Ba 137	Di 138	Ce 140				
9								
10			Er 178	La 180	Ta 182	W 184		Os 195 Ir 197 Pt 198 Au 199
11	(Au 199)	Hg 200	Tl 204	Pb 207	Bi 208			
12				Th 231		U 240		

**C** Mendeleev's 1871 table with his relative atomic masses. The red boxes are gaps left for elements not known at the time. Di, 'didymium', was later shown to be a mixture of two elements, neodymium and praseodymium.

Mendeleev used the gaps in his table to make **predictions** about the properties of undiscovered elements, based on the properties of nearby elements. One set of predictions was for an element he called eka-aluminium. When gallium was discovered shortly afterwards in 1875, its properties closely fitted those Mendeleev had predicted for eka-aluminium.

Property	Eka-aluminium, Ea	Gallium, Ga
relative atomic mass	about 68	70
density of element (g/cm <sup>3</sup> )	about 6.0	5.9
melting point of element (°C)	low	29.8
formula of oxide	Ea <sub>2</sub> O <sub>3</sub>	Ga <sub>2</sub> O <sub>3</sub>
density of oxide (g/cm <sup>3</sup> )	about 5.5	5.88
reacts with acids and alkalis?	yes	yes

**D** Mendeleev's predicted properties of eka-aluminium and the properties of gallium. He also successfully predicted the properties of scandium (discovered in 1879), germanium (1886) and polonium (1898).

### Exam-style question

Give two possible reasons why other scientists did not accept Mendeleev's periodic table when it was first published. (2 marks)

### Did you know?

A 313 km diameter crater on the far side of the Moon was named in honour of Mendeleev by the Russian scientists in charge of the Luna 3 space probe. The probe, launched in 1959, was the first to photograph the far side of the Moon. Element 101 is also named after Mendeleev.



- 3** Mendeleev amended the relative atomic masses of some elements between 1869 and 1871. Give the symbol of one element for which its value was approximately doubled.



- 4** Explain how Mendeleev's 1871 table shows he was unsure where to place three elements.



- 5** Explain why the discovery of gallium was seen as a successful test of Mendeleev's periodic table.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** What were the key features of Mendeleev's periodic table?

### Extend

- E1** How did Mendeleev think creatively to produce his table?  
**E2** What evidence supported Mendeleev's ideas?



# SC4b Atomic number and the periodic table

Specification reference: C1.15; C1.16; C1.17; C1.18

## Progression questions

- Why was Mendeleev right to alter the order of some elements in his table?
- What is an element's atomic number?
- How are the elements arranged in the modern periodic table?



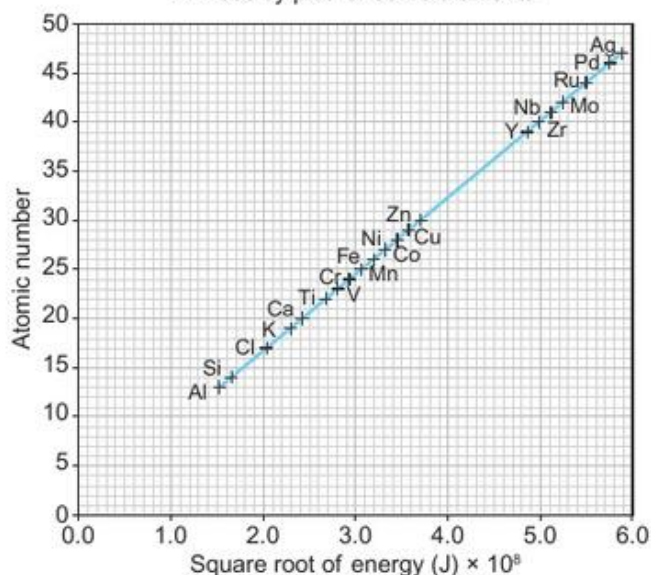
**A** The modern periodic table is easily recognised. There are many fun versions including this one advertising a science park.

Development of the periodic table continued after Mendeleev's first tables. An entire group of **inert** or very unreactive elements was discovered near the end of the 19th century. Even though chemists had not predicted their existence, they were easily fitted into the periodic table as group 0. However, pair reversals such as iodine and tellurium were not properly explained, and there were still gaps. This began to change in 1913 due to a physicist called Henry Moseley.



- 1** Suggest why chemists in Mendeleev's time did not predict the existence of group 0 elements such as neon.

A Moseley plot for some elements



**B** There is a linear relationship between atomic number and the square root of the energy of emitted X-rays.

## Atomic number

When scientists were beginning to accept the periodic table, an element's atomic number was just its position in the table. Moseley showed instead that it is a physical property of an element's atoms. He fired high-energy electrons at different elements, which made them give off **X-rays**. Moseley discovered that for every step increase in atomic number there was a step change in the energy of these X-rays.

Moseley realised that an atomic number was equal to the number of positive charges in the nucleus of an atom. The particle that carries this charge, the proton, was discovered a few years later. So the **atomic number** must be the number of protons in a nucleus.



- 2** Describe the difference between Mendeleev's atomic numbers and Moseley's modern atomic numbers.



## Pair reversals

The elements in the modern periodic table are arranged in order of increasing atomic number,  $Z$ . When this is done:

- elements in a row or **period** are in order of increasing atomic number
- elements with similar properties are in the same column or **group**
- non-metals are on the right of the table (the other elements are metals)
- the iodine–tellurium pair reversal is explained.

Iodine exists naturally as  $^{127}\text{I}$  but tellurium has several different isotopes. About 20% of its atoms are  $^{126}\text{Te}$  but nearly two-thirds of its atoms are  $^{128}\text{Te}$  or  $^{130}\text{Te}$ , so its **relative atomic mass** is greater than that of iodine.

	group 6	group 7	
all (except oxygen) react with oxygen	oxygen, O colourless gas $A_r = 16.0$ $Z = 8$	fluorine, F pale yellow gas $A_r = 19.0$ $Z = 9$	none of them react with oxygen
none of them react with water	sulfur, S yellow solid $A_r = 32.1$ $Z = 16$	chlorine, Cl green-yellow gas $A_r = 35.5$ $Z = 17$	all react with water
all form compounds containing hydrogen: $\text{H}_2\text{O}$ , $\text{H}_2\text{S}$ , $\text{H}_2\text{Se}$ , $\text{H}_2\text{Te}$	selenium, Se metallic grey solid $A_r = 79.0$ $Z = 34$	bromine, Br red-brown liquid $A_r = 79.9$ $Z = 35$	all form compounds containing hydrogen: $\text{HF}$ , $\text{HCl}$ , $\text{HBr}$ , $\text{HI}$
	tellurium, Te silvery-white solid $A_r = 127.6$ $Z = 52$	iodine, I purple-black solid $A_r = 126.9$ $Z = 53$	

**C** These are the elements in groups 6 and 7, each with its relative atomic mass,  $A_r$ , and atomic number,  $Z$ .

## Filling gaps

More X-ray analysis showed that just seven elements between hydrogen ( $Z = 1$ ) and uranium ( $Z = 92$ ) were left to be discovered. These were all discovered between 1917 and 1945. Neptunium, the first element with an atomic number above 92, was discovered in 1940. Other such 'transuranium' elements continue to be discovered, and all can be placed in the periodic table.

- 6** Suggest why there is a gap between calcium, Ca, and titanium, Ti, in graph B.

## Exam-style question

- a** Give an example, other than iodine and tellurium, of a pair of elements that would be in the wrong places if ordered by relative atomic mass. Use a periodic table to help you. (1 mark)
- b** Suggest why ordering by relative atomic mass would be incorrect. (1 mark)

## Did you know?

Not all of Mendeleev's predictions were correct. For example, he predicted an element with an atomic weight of 0.4 that he called coronium. Moseley's work showed that it could not exist because it would need to contain part of a proton.

- 3** Give the relative atomic masses of tellurium and iodine to the nearest whole numbers.
- 4** Use information from diagram C to explain fully why Mendeleev was correct after all to place tellurium before iodine.
- 5** Use the periodic table at the back of the book to find the metals rubidium to tin, and the non-metals in groups 6 and 7. Describe the general positions of metals and non-metals.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** How are the elements arranged in the modern periodic table?

## Extend

**E1** What are the features of the modern periodic table?

# SC4c Electronic configurations and the periodic table

Specification reference: C1.18; C1.19; C1.20

## Progression questions

- What information does an electronic configuration give?
- How do you work out and show the electronic configuration of an element?
- How is the electronic configuration of an element related to its position in the periodic table?

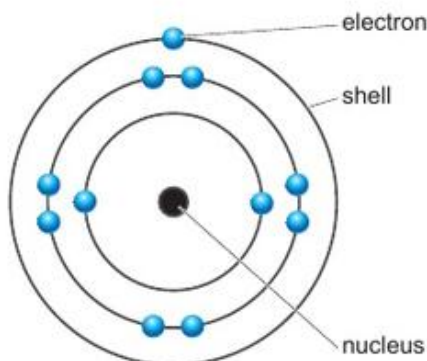


You have many choices where to sit on an empty bus but fewer choices when other people are already seated. **Electrons** fill shells in an atom, rather like filling a bus one seat at a time from the front.

## Electron shells

In an atom, electrons occupy **electron shells** arranged around the nucleus. The shells can be modelled in diagrams as circles, with the electrons drawn as dots or crosses on each circle. The way in which an atom's electrons are arranged is called its **electronic configuration**. Sodium atoms each contain 11 electrons, and diagram B shows the electronic configuration for sodium.

**A** There are many pairs of empty seats on this bus. Where would you sit?



**B** The electronic configuration of sodium shows three occupied shells.

Each shell can contain different numbers of electrons. For the first 20 elements (hydrogen to calcium):

- the first shell can contain up to two electrons
- the second and third shells can contain up to eight electrons.

Electrons occupy the shells, starting with the innermost shell and working outwards as each one becomes full. This is why, in a sodium atom, the first shell contains two electrons, the second shell contains eight electrons and the third shell contains one electron.



**1** State what is meant by the term 'electronic configuration'.



**2** Explain why the electrons in a sodium atom do not all occupy one shell.

## Working out configurations

Electronic configurations can also be written out rather than drawn. For example, the electronic configuration for sodium is 2.8.1 – the numbers show how many electrons occupy a shell, and the full stops separate each shell.

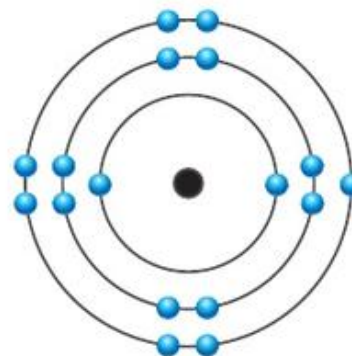
You can work out the electronic configuration of an element from its atomic number, *Z*. The atomic number of chlorine is 17 – each chlorine atom contains 17 protons and so also contains 17 electrons.



To fill a chlorine atom's shells:

- 2 electrons occupy the first shell (leaving 13 electrons)
- 8 electrons occupy the second shell (leaving 5 electrons)
- 5 electrons occupy the third shell.

The electronic configuration of chlorine is therefore 2.8.7 (diagram C shows this).



**C** The electronic configuration of chlorine shows three occupied shells.



- 3** Describe how you can determine the atomic number,  $Z$ , of an element from its electronic configuration.



- 4** Write the electronic configuration of phosphorus,  $Z = 15$ .

## Connections with the periodic table

Diagram D shows the electronic configurations for the first 20 elements in the periodic table. The electronic configuration of an element is related to its position:

- the number of occupied shells is equal to the period number
- the number of electrons in the outer shell is equal to the group number (except for group 0 elements, which all have full outer shells).

1	2	3	4	5	6	7	0

**D** These are the electronic configurations of the first 20 elements.



- 5** What do the electronic configurations of sodium and the other elements in group 1 have in common?



- 6** Explain how you can tell from their electronic configurations that sodium and chlorine are in the same period.

## Exam-style question

Explain, in terms of electrons, why magnesium and calcium are in the same group in the periodic table. (2 marks)

## Did you know?

After calcium, the third and fourth electron shells can actually contain up to 18 electrons. For this reason the International Union of Pure and Applied Chemistry (IUPAC) now recommends the use of group numbers in the range 1 to 18. In this newer numbering system, group numbers 3 to 12 refer to the block of elements between calcium and gallium. Group numbers 13 to 17 refer to the older group numbers 3 to 7, and 18 refers to group 0.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** How do you work out the electronic configuration of an element?

## Extend

- E1** How is the electronic configuration of an element determined, and related to its position in the periodic table?

# Atomic structure and the periodic table

The table shows information about tellurium and iodine.

Element	Atomic number	Relative atomic mass, $A_r$
tellurium	52	127.6
iodine	53	126.9

**A**

Describe how Mendeleev arranged the elements known to him into a periodic table, and compare this with the arrangement in the modern periodic table.

In your answer, use information from the table above to provide examples.

(6 marks)

## Student answer

Mendeleev arranged the elements in order of relative atomic mass [1]. He also used the properties of elements and their compounds, so he sometimes had to swap elements. For example, the  $A_r$  of iodine is 126.9, so he should have put iodine first [2], but he put it second. He did this so it went into the same column as similar elements [3].

In the modern periodic table, elements are arranged in order of increasing atomic number (the number of protons in the nucleus) [4]. This explains why Mendeleev was correct when he placed tellurium before iodine.

[1] This shows knowledge of how Mendeleev ordered the elements, but it should say 'in order of increasing relative atomic mass'.

[2] The answer should be supported by comparing the  $A_r$  values for the two elements.

[3] An element in the same group, such as chlorine, could be identified using the periodic table.

[4] It is a good idea to give clear definitions like this.

## Verdict

This is an acceptable answer. It shows good knowledge and understanding of the periodic table and the answer is organised in a logical way. The student has linked their knowledge of atomic structure with an understanding of Mendeleev's periodic table.

The answer could be improved by clearly quoting examples from the information given in the table. For example, in the last paragraph the answer could have included the atomic numbers for each of the elements.

### Exam tip

If you are asked to use information from the question as part of your answer then make sure you include the information. It is important to make sure you *use* the information in your answer – don't just repeat what you have been given.



## Paper 1 and Paper 2

### SC5 Ionic Bonding / SC6 Covalent Bonding / SC7 Types of Substance

Using an extremely powerful 'atomic force microscope' scientists at the Berkeley Lab in California produced this amazing image showing the positions of the bonds holding the atoms of a small molecule together. Using this technique, the scientists were also able to examine the breaking and reforming of bonds during a chemical reaction. Bonds are the fundamental forces of attraction that hold our universe together. Understanding how bonds are formed and broken is essential in helping us explain even the simplest physical change or chemical reaction.

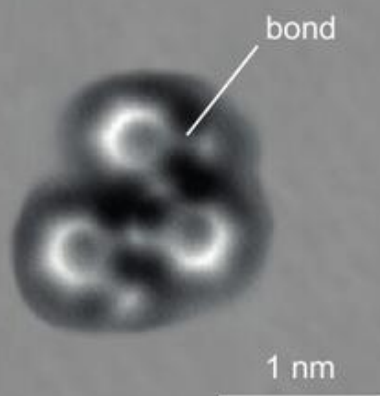
#### The learning journey

Previously you will have learnt at KS3:

- about the particle model of matter
- how Dalton's ideas about atoms and molecules helped to explain the properties of matter
- how elements are arranged in the periodic table.

In this unit you will learn:

- how ionic, covalent and metallic bonds are formed
- about the formation of lattice and molecular structures
- how the physical properties of a substance are linked to its bonding and structure.

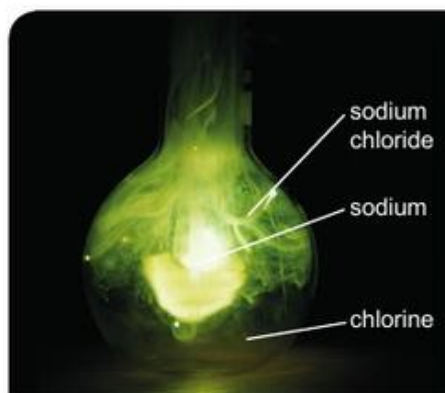


# SC5a Ionic bonds

Specification reference: C0.1; C1.21; C1.22; C1.23; C1.24

## Progression questions

- How are ions formed?
- How can the numbers of subatomic particles in an ion be calculated?
- What is an ionic bond?

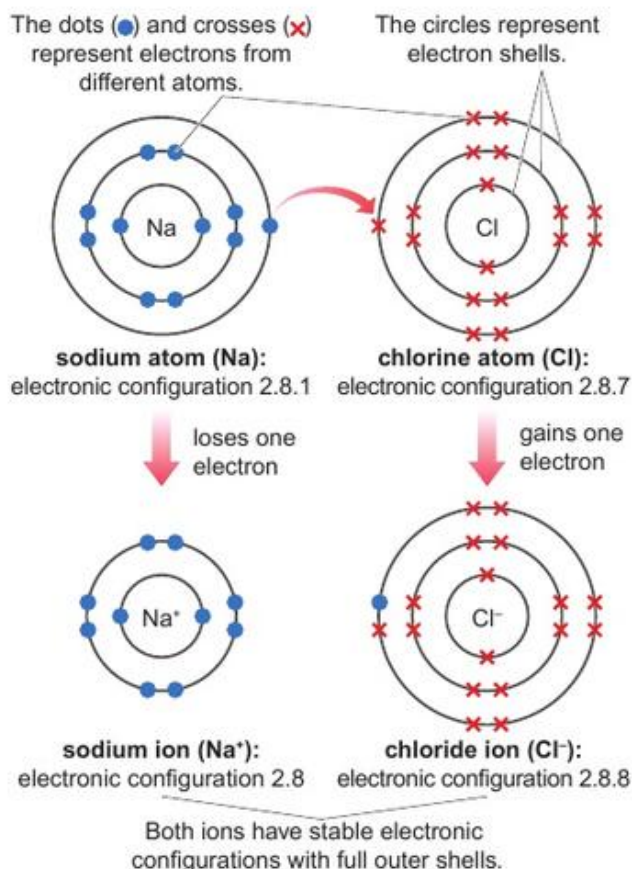


**A** Explosions can occur when new bonds form between sodium and chlorine atoms to form sodium chloride.

**Bonds** are forces of attraction that hold atoms together. When bonds form between atoms, energy is released from the atoms, making them more stable (less reactive). The most stable atoms are those of the noble gases, and scientists have found that this is because they have outer electron shells that contain as many electrons as possible – the outer shell of a noble gas atom is full.

Noble gas	Electronic configuration
He	2
Ne	2.8
Ar	2.8.8

**B** Atoms of all of the elements in group 0 have a stable electronic configuration (arrangement of electrons) with a complete outer shell of electrons.



**C** Dot and cross diagrams can be used to show what happens when ions are formed.

Atoms are more stable if they have an outer electron shell that is full, like a noble gas. This can happen by the transfer of electrons between atoms, forming charged particles called **ions**.

Metal atoms tend to lose electrons and form positive ions, called **cations**. Cations have more protons than electrons. Non-metal atoms tend to gain electrons and form negative ions, called **anions**. Anions have more electrons than protons. The formation of sodium (Na<sup>+</sup>) and chloride (Cl<sup>-</sup>) ions is shown in diagram C. Note that when non-metals form negative ions the end of the name changes to -ide.



**1 a** What are ions?



**b** State the difference between a cation and an anion.

There are forces of attraction between all positively and negatively charged objects. These are called **electrostatic forces**. These forces hold the oppositely charged ions together, and form an **ionic bond** between them.



**2 a** What is happening to the atoms when they form ions?



**b** What holds the ions together in an ionic bond?

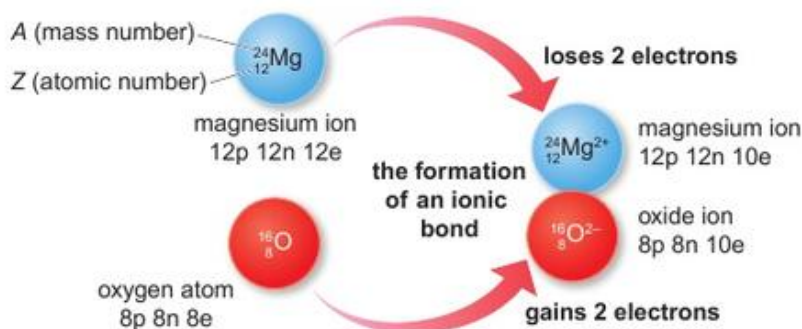


Atoms that easily form ions will have either a nearly full or a nearly empty outer electron shell. Most ionic bonds are formed between a metal and a non-metal. Table D shows the number of electrons lost or gained and the resulting ion charge of some groups of elements. The ion formed depends on the element's position in the periodic table and its number of outer electrons.

	Group 1	Group 2	Group 6	Group 7
outer electrons	1	2	6	7
electrons lost or gained	1 lost	2 lost	2 gained	1 gained
charge on ion	1+	2+	2-	1-
example	Li <sup>+</sup>	Ca <sup>2+</sup>	S <sup>2-</sup>	F <sup>-</sup>

**D** formation of ions by elements in groups 1, 2, 6 and 7

Diagram E shows the number of protons, neutrons and electrons in each atom and ion in the formation of the ionic bond in magnesium oxide.



**E** ionic bond formation in magnesium oxide

- 5 An aluminium atom  $^{27}_{13}\text{Al}$  loses three electrons when its ion is formed.



**a** How many protons, neutrons and electrons are in the aluminium ion?



**b** Write the symbol for the aluminium ion.



- 6 Draw dot and cross diagrams to show the ions in diagram E.

### Exam-style question

Bromine is in group 7 of the periodic table.

- a** Write the symbol for a bromide ion. (1 mark)
- b** State what the symbol tells you about the electrons in a bromide ion compared with a bromine atom. (1 mark)

### Did you know?

The most common atom in the Universe, hydrogen, contains one proton and one electron. The hydrogen ion ( $\text{H}^+$ ) is therefore just a proton. This makes it more than 100 000 times smaller than the hydrogen atom and it is the smallest particle that can take part in a chemical reaction.

- 3 Write the symbol for the ion formed by:



**a** potassium (K) in group 1



**b** selenium (Se) in group 6.

- 4 A sulfur atom contains 16 protons.



**a** How many electrons does it contain?



**b** Write out its electronic configuration.



**c** Explain why it forms  $\text{S}^{2-}$  ions.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Describe what happens when lithium (group 1) and fluorine (group 7) react to form an ionic bond.

### Extend

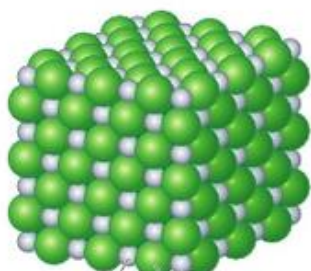
**E1** Using dot and cross diagrams, explain what happens when aluminium and oxygen form an ionic bond.

# SC5b Ionic lattices

Specification reference: C0.1, C1.25, C1.26, C1.27

## Progression questions

- What is an ionic lattice?
- What do the endings -ide and -ate tell you about a substance?
- How do you work out the formulae of ionic compounds?



Na<sup>+</sup> (a sodium ion)    Cl<sup>-</sup> (a chloride ion)

**A** Sodium chloride forms a cubic lattice structure.



- 2** Explain why the solid shown in photo B can be described as crystals.



**B** Sodium chloride crystals are shaped like cubes. These crystals are from a salt mine in Poland and the largest one has sides of 5 cm.

**Ionic compounds**, which are formed by the loss and gain of electrons, are held together by strong electrostatic forces of attraction between oppositely charged ions. These strong ionic bonds allow 'billions' of ions to be packed together in a regular repeating arrangement called a **lattice structure**. The lattice structure in sodium chloride is shown in diagram A.



- 1 a** What holds ions together in an ionic bond?

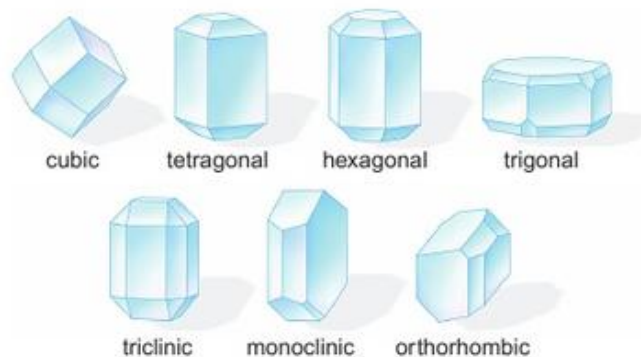


- b** Why is the structure in diagram A described as a cubic lattice?

Ionic compounds will often form **crystals** when solid because of their regular lattice structure. Crystals are pieces of solid that have a particular regular shape, flat surfaces and sharp edges. Photo B shows crystals of sodium chloride.

## Did you know?

The shapes of crystals are determined by the structure of the lattice. There are seven basic crystal or lattice structures.



**C** the seven basic crystal structures

Positive ion	Ion formula	Negative ion	Ion formula
sodium	Na <sup>+</sup>	fluoride	F <sup>-</sup>
lithium	Li <sup>+</sup>	chloride	Cl <sup>-</sup>
potassium	K <sup>+</sup>	bromide	Br <sup>-</sup>
magnesium	Mg <sup>2+</sup>	oxide	O <sup>2-</sup>
calcium	Ca <sup>2+</sup>	sulfide	S <sup>2-</sup>
aluminium	Al <sup>3+</sup>	phosphide	P <sup>3-</sup>

**D** common ions

## Working out ionic formulae

Ionic compounds are electrically neutral (they have no overall charge). So the formula of an ionic compound contains the same number of positive charges as negative charges. To work out ionic formulae we will need to use ion formulae, as shown in table D. Note that in two-element compounds the name ending of the non-metal ion is changed to -ide.



**Worked example W1**

Magnesium oxide contains the ions  $\text{Mg}^{2+}$  and  $\text{O}^{2-}$ .

A magnesium ion has two positive charges and an oxide ion has the same number of negative charges. One of each ion will balance the charges, and so the formula =  $\text{MgO}$

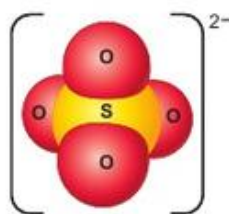
**Worked example W2**

Sodium sulfide contains the ions  $\text{Na}^+$  and  $\text{S}^{2-}$ .

Therefore two  $\text{Na}^+$  ions are needed to balance the charges on the  $\text{S}^{2-}$  ion, and so the formula =  $\text{Na}_2\text{S}$

Some ions contain more than one atom. For example, the sulfate ion, shown in diagram E, contains one sulfur atom bonded to four oxygen atoms with two added electrons. More examples of these **polyatomic ions** are shown in table F. An ion name ending '-ate' or '-ite' shows that the ion contains oxygen as well as another element.

If an ionic formula contains two or more of the same polyatomic ions then the formula of the polyatomic ion must be written inside brackets.



**E** The formula of a sulfate ion is  $\text{SO}_4^{2-}$ .

**Worked example W3**

Calcium nitrate contains the ions  $\text{Ca}^{2+}$  and  $\text{NO}_3^-$ .

Therefore two  $\text{NO}_3^-$  ions are needed to balance the charges on the  $\text{Ca}^{2+}$  ion, and so the formula =  $\text{Ca}(\text{NO}_3)_2$

Note that the brackets are needed around the polyatomic ion. Without brackets the formula becomes  $\text{CaNO}_{32}$  and this compound is not possible.

**5** Write the formulae for:



**a** sodium carbonate

**b** ammonium sulfate.

**6**  $\text{Li}_3\text{PO}_4$  is an ionic compound.



**a** Suggest a name for this compound.



**b** Write the formulae for both of the ions.

**Exam-style question**

Potassium iodide and potassium iodate, which contains the ion  $\text{IO}_3^-$ , are both used in health supplements.

**a** Give the formulae for both compounds. (1 mark)

**b** Compare and contrast these two compounds. (2 marks)

**3** Write the formulae for:



**a** sodium fluoride



**b** calcium sulfide



**c** magnesium bromide



**d** aluminium oxide.

**4** The formula of iron chloride is  $\text{FeCl}_3$ .



**a** What is the charge on the iron ion?



**b** Explain how you worked out your answer.

Polyatomic ion name	Ion formula
ammonium	$\text{NH}_4^+$
nitrate	$\text{NO}_3^-$
hydroxide	$\text{OH}^-$
carbonate	$\text{CO}_3^{2-}$
sulfate	$\text{SO}_4^{2-}$
sulfite	$\text{SO}_3^{2-}$

**F** Polyatomic ions are groups of two or more atoms, which have become charged.

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

**S1** Describe with the help of a diagram the type of structure you find in ionic compounds.

**Extend**

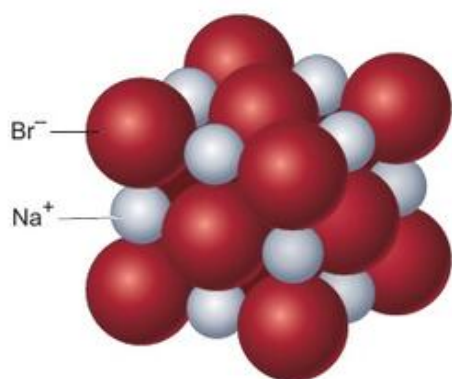
**E1** What does the formula  $\text{Fe}_3(\text{PO}_4)_2$  tell you about the compound and its structure?

# SC5c Properties of ionic compounds

Specification reference: C1.33

## Progression questions

- What particles and forces are present in ionic compounds?
- Why do ionic compounds have high melting points and boiling points?
- Why do ionic compounds conduct electricity when they are liquids or dissolved in water but not when they are solids?



A the lattice structure of sodium bromide

All ionic compounds contain charged particles called ions. Ions have one or more positive charges or one or more negative charges. The oppositely charged ions in ionic compounds are held together by strong electrostatic forces of attraction, which we call ionic bonds. It is important to remember that ionic compounds contain *ions*, not atoms or molecules.

In ionic compounds, huge numbers of ions are arranged in a giant structure or lattice. This has a regular pattern and so ionic compounds form crystals.



1 State the type of particles present in an ionic compound.



2 State the type of forces found in an ionic bond.



B molten sodium chloride

## Melting points and boiling points

The electrostatic forces of attraction between oppositely charged ions are strong. A lot of energy is needed to overcome these forces in order to separate the ions and cause the substance to melt. This is why ionic compounds must be heated to high temperatures before they change state. They have high melting points and boiling points.

Some ions have more than one charge (such as  $\text{Mg}^{2+}$  and  $\text{O}^{2-}$ ). These highly charged ions will attract other ions more strongly than ions with one charge. More energy will be needed to overcome the electrostatic forces of attraction and so the melting points will be higher.

Ionic compound	Melting point ( $^{\circ}\text{C}$ )	Boiling point ( $^{\circ}\text{C}$ )
sodium bromide, NaBr	747	1390
sodium chloride, NaCl	801	1413
magnesium oxide, MgO	2852	3600

C Ionic compounds have high melting points and boiling points.



3 Explain why ionic compounds have high melting points.



4 Explain why magnesium oxide has a much higher melting point than sodium chloride.

## Electrical conductivity

Ionic compounds conduct electricity when they are molten or dissolved in water. They do not conduct electricity when they are in the solid state.

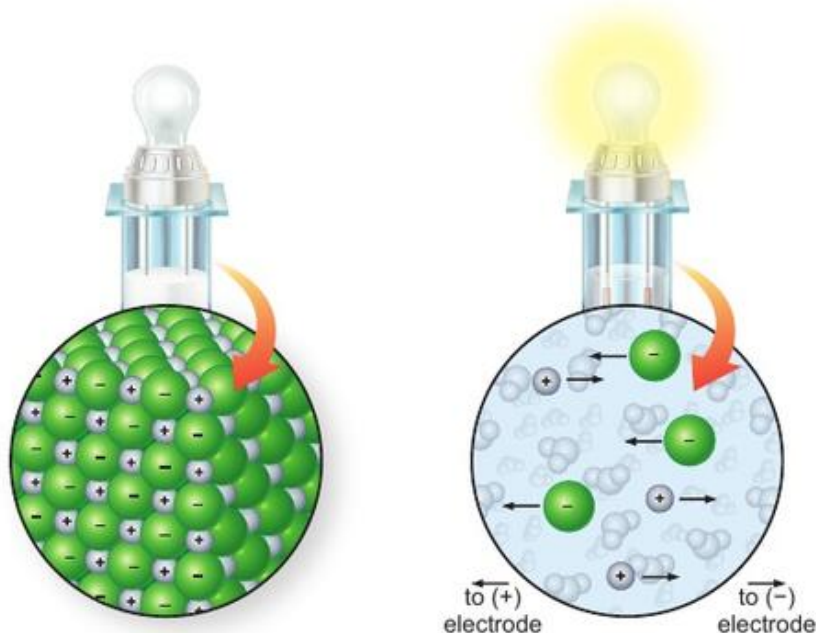
Two conditions must be met for a substance to conduct electricity:

- it must contain charged particles
- these particles must be free to move.



- 5** Samples of sodium chloride and magnesium oxide are both heated to  $1080^{\circ}\text{C}$ . State and explain how the properties of these two compounds will or will not change at this temperature compared to room temperature.

When an ionic compound conducts electricity, it is the charged *ions* that carry the current. Ionic compounds do not conduct electricity in the solid state as the ions are not free to move from place to place. When the ionic compound is molten or in **aqueous solution**, the ions are free to move and so it does conduct electricity. Most ionic compounds are soluble in water and form an aqueous solution.



Positive and negative ions fixed in a solid do not conduct a current.

In solution, positive and negative ions move and conduct a current.

**D** Solid sodium chloride does not conduct electricity but aqueous sodium chloride does conduct.

The negative ions are also known as anions and they are attracted to the positive electrode, which is called the **anode**. The positive ions are also known as cations and they are attracted to the negative electrode, which is called the **cathode**.

It is important to remember that it is ions moving that enable ionic compounds to conduct electricity and that it is not *electrons* moving.

- 6** Explain why ionic compounds conduct electricity when they are dissolved in water.
- 7** Magnesium oxide is insoluble in water. State the condition needed for magnesium oxide to conduct electricity.

### Exam-style question

Magnesium chloride is an ionic compound and has a high melting point. Explain why magnesium chloride has a high melting point. (3 marks)

### Did you know?

Molten salts are used to store energy in some solar power stations. The salts are heated up by the Sun and release their thermal energy at night, allowing electricity to be generated when it is dark.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** You are given a substance and asked to find out if it is an ionic compound. Describe the tests you would carry out in order to decide.

### Extend

**E1** Table E shows some properties of five compounds.

Compound	Melting point ( $^{\circ}\text{C}$ )	Soluble in water?	Conducts when molten?
A	2072	no	yes
B	191	no	no
C	782	yes	yes
D	605	yes	yes
E	150	yes	no

### E

Identify the compounds that have ionic bonding and explain your reasoning. Give reasons why the other compounds do not have ionic bonding.

# SC6a Covalent bonds

Specification reference: C1.28; C1.29; C.1.30; C1.31

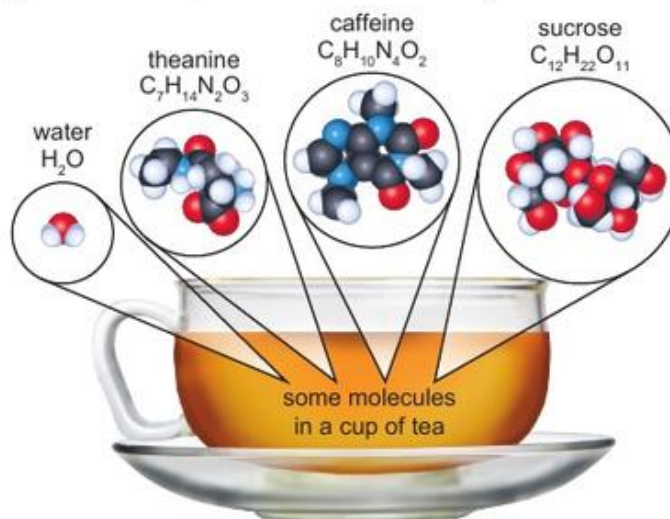
## Progression questions

- What are the names of some covalent molecules?
- How are covalent bonds formed?
- How can dot and cross diagrams be used to explain the formation of covalent molecules?

### Did you know?

A cup of tea will contain over 1 000 000 000 000 000 000 000 (10<sup>24</sup>) water molecules. This is the same as the estimated number of stars in the Universe.

**Molecular** substances contain groups of atoms that are held together by strong bonds called **covalent bonds**. The number of atoms of each element bonded together in a simple **molecule** is shown by its **molecular formula**.



- What is a molecule?
  - What does a molecular formula tell you about a substance?

- Oxygen molecules consist of two oxygen atoms.

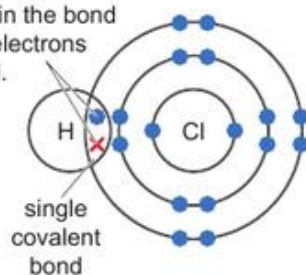
- Write the molecular formula for oxygen.
- Explain why a noble gas, such as argon, exists as single atoms and not as molecules.

Dots show electrons from one atom and crosses show electrons from the other atom. This allows us to see which atoms the electrons in the bond originally came from. The electrons themselves are all identical.

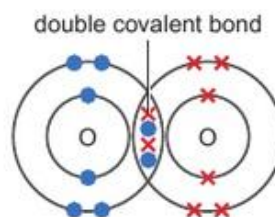
single covalent bond



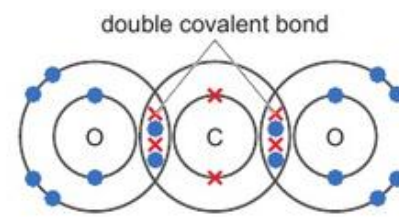
a hydrogen



b hydrogen chloride



c oxygen



d carbon dioxide

**B** Dot and cross diagrams can be used to explain how covalent bonds are formed.

**A** Most of the molecules in a cup of tea are water, which has the molecular formula H<sub>2</sub>O.

Covalent bonds are usually formed between non-metal atoms and are produced by sharing pairs of electrons. By forming the bond the atoms become more stable, because they can use the shared electrons to complete their **outer electron shells**. The reason why noble gases are so stable is because they have full outer electron shells.

The **dot and cross diagrams** in diagram B show how covalent bonds are formed. Counting the shared electrons, each atom now has a complete outer shell of electrons. Sometimes atoms share more than one pair of electrons to fill their outer shells. In oxygen and carbon dioxide the atoms share two pairs of electrons, to form **double bonds**.





- 3** a Describe how covalent bonds are formed.  
 b Write the molecular formulae for the substances shown in diagram B.
- 4** The electronic configuration of fluorine is 2.7. Draw a dot and cross diagram to show how fluorine molecules ( $F_2$ ) are formed.

The atoms in molecules are held together by strong **electrostatic forces** of attraction between the positive nuclei and the negative electrons in the bonded atoms. There are also some forces of attraction *between* molecules but these are very weak in comparison. Atoms and molecules are extremely small, about  $10^{-10}$  metres across, so we represent them using models, such as those shown in Diagram C.

- 5** What holds the atoms together when a covalent bond has been formed?

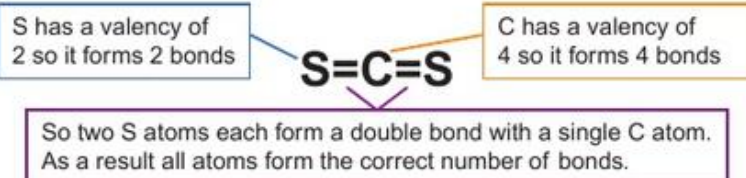
### Working out molecular formulae

The numbers of covalent bonds formed by atoms of different elements are shown in table D. This is called the **valency** of the element. It is the same as the number of electrons needed to obtain a complete outer shell.

Group number	Examples	Outer electrons	Bonds formed	Valency
4	C and Si	4	4	4
5	N and P	5	3	3
6	O and S	6	2	2
7	F and Cl	7	1	1

#### D valencies of some elements

Diagram E shows how molecular formulae can be worked out by matching up the valencies, so that all atoms have the correct number of bonds (and so a complete outer electron shell).



#### E working out the formula of carbon sulfide

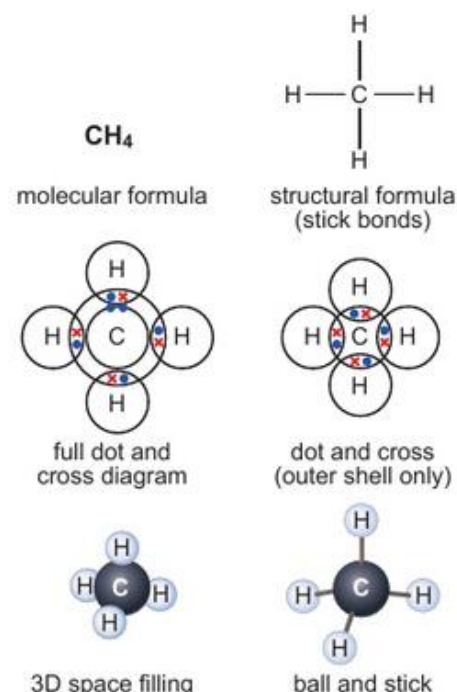
**7** Work out the molecular formulae for:

- a oxygen fluoride    b hydrogen sulfide    c nitrogen chloride.

### Exam-style question

Describe the bonding in a molecule of water, which contains two hydrogen atoms and one oxygen atom.

(3 marks)



#### C different representations of methane

**6** Look at diagram C.

- a How are bonds represented in structural formulae?
- b Describe the covalent bonds in methane.
- c Draw four different models of a water molecule, including one dot and cross diagram.

### Checkpoint

How confidently can you answer the Progression questions?

#### Strengthen

**S1** What does diagram B tell you about hydrogen chloride?

#### Extend

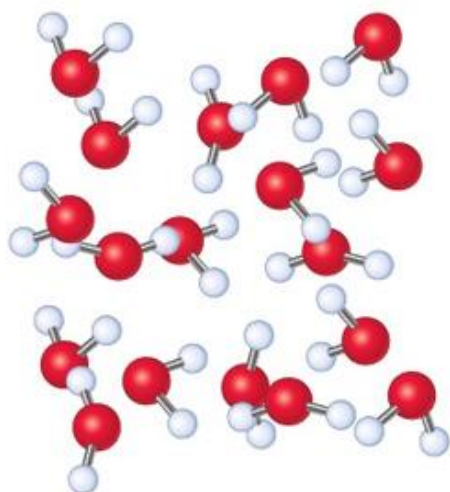
**E1** Describe the bonding in a molecule of ammonia, which contains three hydrogen atoms and one nitrogen atom. Use diagrams in your answer.

# SC7a Molecular compounds

Specification reference: C1.34; C1.39

## Progression questions

- Why do simple molecular compounds have low boiling and melting points?
- Why are simple molecular compounds poor electrical conductors?
- What is a polymer?



A 'ball and stick' models of  $\text{H}_2\text{O}$  molecules

All **compounds** contain atoms of more than one **element**, chemically joined together by **bonds**. The properties of a compound are influenced by its atoms and by its type of bonding.

Some compounds exist as molecules – distinct groups of atoms joined by **covalent bonds**. They have **covalent, simple molecular structures**. An example is water. One molecule of water always contains one atom of oxygen covalently bonded to two atoms of hydrogen.

1 Look at diagram A.



a What do the red and white spheres represent?



b How are bonds represented in this model?



2 Why is the ratio of oxygen to hydrogen atoms the same in every glass of water?

## Did you know?

Intermolecular forces can allow water in an open container to be higher than the top of the container.



B

## Melting and boiling points

The covalent bonds in a water molecule are strong forces of attraction. However, there are also weak forces of attraction *between* molecules – **intermolecular forces**. These intermolecular forces hold water molecules together and must be overcome when turning liquid water into a gas. Small, simple molecules such as water often have low melting and boiling points, because it doesn't take much energy to overcome the weak intermolecular forces.



3 Covalent bonds are strong, so why does water have a low melting point?



4 What would you expect the strength of the bonds to be like between carbon and oxygen atoms in carbon dioxide, compared to the strength of the forces between neighbouring carbon dioxide molecules?



5 a Methane consists of molecules (containing one carbon atom covalently bonded to four hydrogen atoms). Would you expect methane to have a high or low melting point?



b Explain your reasoning.



## Conduction of electricity

An electric current is a flow of charged particles. Simple molecules have no overall charge and so cannot carry an electric current. In a covalent bond, electrons are shared between two atoms. The strong forces between the negatively charged electrons and the positively charged nuclei hold the electrons in place. The electrons cannot flow and so cannot carry a current.



- 6** Explain why water is a poor conductor of electricity.

## Polymers

**Monomers** are small, simple molecules that can be joined in a chain to form a **polymer**. Carbon atoms can form up to four covalent bonds with other atoms, and so monomers are usually linked together by covalent bonds between carbon atoms. Most polymers contain a chain of carbon atoms.

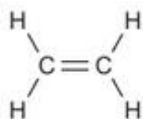
**Poly(ethene)** (or 'polythene') is a common polymer made of ethene monomers (as shown in diagram D).

Polymer molecules can have different lengths. Longer polymers have more intermolecular forces between them. The longer chains also tend to get tangled up with one another. For these reasons, longer polymers have higher melting and boiling points than shorter ones.

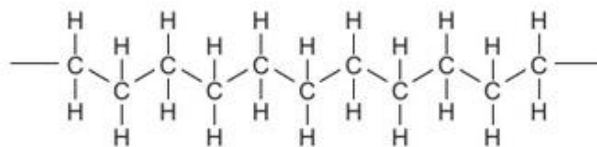


**C** Poly(ethene) sheets are often used to protect things, such as plants. These are coffee beans being dried.

(i) ethene



(ii) poly(ethene)



**D** two dimensional models of: (i) an ethene molecule (monomer)

(ii) a poly(ethene) molecule



- 8 a** What monomer is used to make poly(butene)?



- b** What polymer is formed by linking styrene monomers together?



- 9** Explain why most polymers are solids at room temperature, whereas their monomers are liquids or gases.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** You can ask three questions to find out if a substance consists of simple molecules or not. What questions would you ask?

### Extend

- E1** Propene melts at  $-185^{\circ}\text{C}$ . Explain how the melting point and electrical conduction will change if other propene molecules are added together to form a chain.

### Exam-style question

Explain why small molecules have a lower melting point than large polymer molecules.

(3 marks)

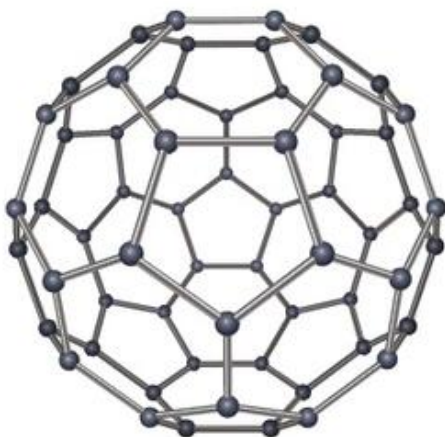
# SC7b Allotropes of carbon

Specification reference: C1.35; C1.36; C1.37; C1.38

## Progression questions

- How are simple molecular structures different from giant covalent structures?
- What are the differences in structure between the different allotropes of carbon?
- How do we explain the properties and uses of graphite, diamond and fullerenes?

### 5<sup>th</sup> 1 What is an allotrope?



**A** Buckminsterfullerene is a simple molecule.

- 6<sup>th</sup> 2 A fullerene has the formula  $C_{60}$ . Describe its structure.
- 6<sup>th</sup> 3 Explain why fullerenes have low melting points.

### Did you know?

Buckminsterfullerene is named after an American architect called Richard Buckminster Fuller, who designed spherical buildings with a similar structure to  $C_{60}$ .

- 6<sup>th</sup> 4 Explain why graphene is not a simple molecule.

Molecules are groups of atoms joined by covalent bonds. Molecules can be compounds (such as water,  $H_2O$ ) or elements (such as oxygen,  $O_2$ ).

The element carbon can form a number of different molecules. Different structural forms of the same element are called **allotropes**. The structure and bonding in different allotropes influences their properties and uses.

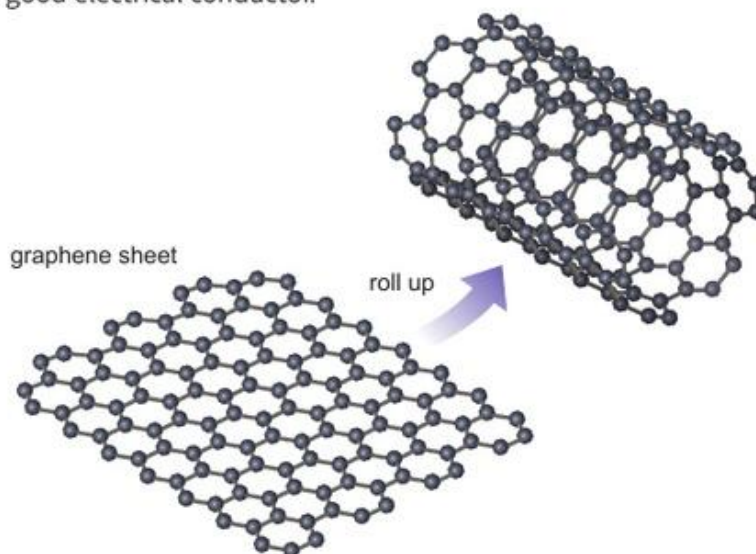
## Fullerenes

Carbon can form simple molecules called **fullerenes**, in which each carbon atom is covalently bonded to three other carbon atoms. Fullerenes are often tubular molecules (nanotubes) or spherical. An example is buckminsterfullerene (or 'bucky ball'), which has 60 carbon atoms that form a ball with the formula  $C_{60}$ .

Fullerenes have weak intermolecular forces between the molecules and so have low melting points (or sublimation points). These weak forces also make them soft and slippery. However, the molecules themselves are very strong due to their covalent bonding.

## Graphene

**Graphene** is similar to fullerenes but is not a simple molecule. It consists of a sheet of carbon atoms with no fixed formula. The sheet is just one atom thick, making it the lightest known material, but its covalent bonds make it extremely strong. It also allows free electrons to move across its surface and so is a good electrical conductor.



**B** Graphene can be a sheet or can be rolled into a tube.



## Giant structures of carbon

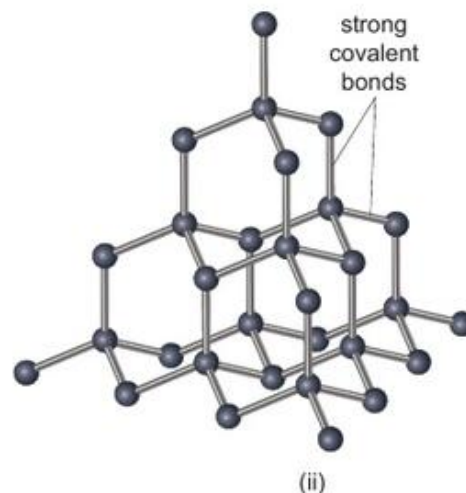
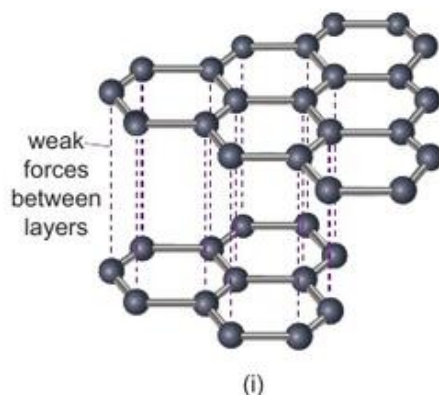
Diamond and graphite are two more allotropes of carbon that are not simple molecules. They are both examples of **covalent, giant molecular structures**, which have huge three-dimensional networks of carbon atoms linked by covalent bonds.



- 5** How many covalent bonds will most carbon atoms form in graphite, diamond and  $C_{60}$ ?



- 6** Describe how simple molecular structures are different from giant molecular structures.



**C** small sections of (i) graphite and (ii) diamond to show the arrangements of atoms

Graphite and diamond both have high melting points because of the many strong covalent bonds that need to be broken to melt the solids.

However, graphite has three covalent bonds for each carbon atom, whereas diamond has four. This gives graphite a layered structure and means that not all of its electrons are held in covalent bonds. These **delocalised electrons** are free to move and can carry an electrical current. Since graphite conducts electricity well and is cheap and not very reactive, it is used as electrodes (in electrolysis).



- 7** Explain why graphite conducts electricity but diamond does not.

The sheets of carbon atoms in graphite are held together by weak forces of attraction (purple dashed lines in diagram C). These weak forces allow the layers to slide past each other, which makes graphite quite soft and useful as a **lubricant**.

Diamond is very hard because it has a rigid network of carbon atoms in a tetrahedral arrangement, joined by strong covalent bonds. This property makes diamond useful for tools to cut things. It is also an electrical insulator because it has no free charged particles.

- 8** Explain why:



**a** graphite is used as electrodes



**b** diamonds are used on cutting heads when drilling through rocks



**c** spherical fullerenes are used in some lubricants.



**D** This cutting head, used in mining, is studded with diamonds.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Describe how the structures of four allotropes of carbon cause their properties.

### Extend

**E1** Graphene sheets can be rolled into tubes. Predict the properties of these tubes.

### Exam-style question

Explain why graphite is softer than diamond.

(4 marks)

# SC7c Properties of metals

Specification reference: C1.40; C1.42

## Progression questions

- What are the typical physical properties of metals and non-metals?
- How are the particles arranged in metals?
- How can we explain the properties of a metal in terms of its bonding and structure?



The bucket is made of iron. Metals have some common properties:

- solids with high melting points
- shiny (when polished)
- malleable
- high density
- good conductors of electricity.

Sulfur is a non-metal. Non-metals have some common properties:

- solids, liquids or gases with low melting points
- not usually shiny (when solid)
- brittle (when solid)
- low density
- poor conductors of electricity.

Chemists classify elements into **metals** and **non-metals**, depending on their properties. Just over three-quarters of the elements are metals.

A metal or a non-metal may not have all the common properties shown in photo A.

1 Give two general properties of:



a metals



b non-metals.

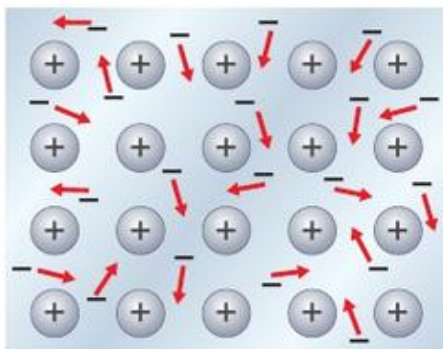


2 Name a form of the non-metal carbon that is a good conductor of electricity.

A Sulfur being unloaded at a port in Darwin, Australia.

## Metallic structure and bonding

The atoms in a metallic element are all the same size and are packed closely together in layers to form a giant **lattice**.



B Metals consist of stacked layers of ions in a 'sea' of delocalised ('free') electrons.

Metal atoms have one, two or three electrons in their outer shell. These outer shell electrons are lost from each atom and become free to move randomly throughout the metal. This leaves a giant lattice of positive metal ions surrounded by a 'sea' of delocalised electrons, which move randomly in all directions.

**Metallic bonding** is the electrostatic attraction between the positive metal ions and the negative delocalised electrons. This attraction is strong, so metals have high melting and boiling points.



3 a Describe the structure of a metal.

b Describe the bonding in a metal.

4 Explain why most metals have high melting points.

## Did you know?

Osmium is the most dense metal at room temperature (the mass of 1 cm<sup>3</sup> is 22.6 g). A lump of osmium the size of an average smartphone has a mass of 1.5 kg (the same as one and a half bags of sugar)!



## Metals are malleable

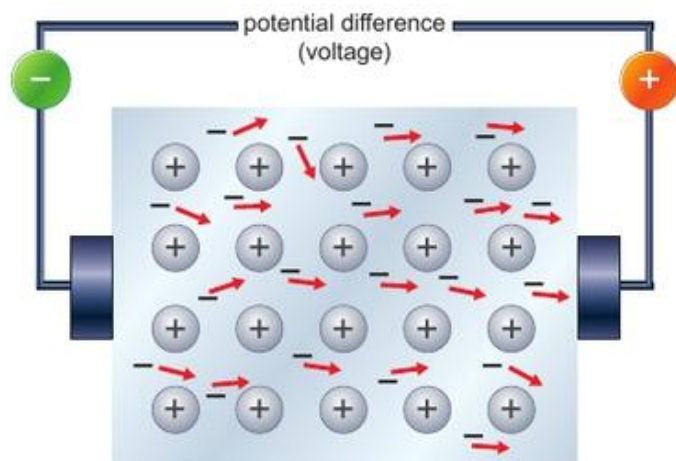
Metals are **malleable**. This means that they can be hammered or rolled into shape without shattering. When you hit a metal, the layers of ions slide over each other. The 'sea' of electrons holds the ions together and so the metal changes shape instead of breaking.



- 5 Copper is used to make some saucepans. Explain why copper can be bent and shaped into a saucepan.

## Metals conduct electricity

The delocalised electrons move randomly between the positive metal ions in all directions. When a potential difference (voltage) is applied between two points on a piece of metal, the electrons will flow towards the positive side. This flow of electrons transfers energy and forms an electrical current.



- E When a voltage is applied to a piece of metal, an electrical current flows.

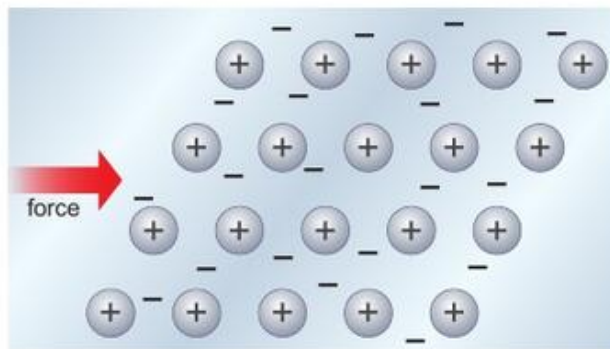


- 6 Copper is used to make electrical wires. Explain why copper is a good conductor of electricity.

Substances that conduct electricity better than others have a higher **electrical conductivity**. The electrical conductivity of the metals increases as the number of delocalised electrons increases. Each sodium ion has one positive charge,  $\text{Na}^+$ , and contributes 1 electron to the 'sea' of delocalised electrons. Each magnesium ion has two positive charges,  $\text{Mg}^{2+}$ , and contributes 2 electrons to the 'sea' of delocalised electrons, and so magnesium has a higher electrical conductivity than sodium.



- 7 Aluminium forms  $\text{Al}^{3+}$  ions. Explain why aluminium has a higher electrical conductivity than magnesium.



- C When hit or bent, the layers of ions in a metal can slide over each other.



- D Copper conducts electricity well and so is used for electrical wiring.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1 You are given a solid substance and asked to find out if it is a metal or a non-metal. Describe the properties you would need to find out about.

### Extend

- E1 Elements A and B are solids at room temperature. Element A has a melting point of  $98^\circ\text{C}$ , a density of  $0.97\text{ g/cm}^3$  and conducts electricity. Element B has a melting point of  $217^\circ\text{C}$ , a density of  $4.81\text{ g/cm}^3$  and doesn't conduct electricity. Are these elements metals or non-metals? Explain your reasoning.

### Exam-style question

Explain how metals conduct electricity.

(2 marks)



# SC7d Bonding models

Specification reference: C1.32; C1.41

## Progression questions

- What different types of structure and bonding models are used to describe substances?
- How do these models help explain the properties of substances?
- What are the limitations of the models that we use to show structure and bonding?



A 3D models of some gas molecules

- 1 Look at the substances in the box below.

potassium fluoride  
carbon dioxide  
diamond  
magnesium

Which substance:



- a conducts electricity when solid



- b is most likely to dissolve in water



- c has the lowest melting point?



- 2 a Name the type of bonding and structure that usually has a low melting point.



- b Explain why the melting points of these substances are usually low.

Scientists have developed models to explain how different types of bonds and structures are formed. These models help us explain the properties of different substances. Most elements and compounds fit into one of the four main models summarised below.

### Ionic

Where found: in most compounds containing metal and non-metal atoms.

Bonding: ionic bonds formed by the loss and gain of electrons to produce oppositely charged ions that attract one another.

Structure: billions of ions held together in a lattice structure.

Properties:

- high melting/boiling points
- many are soluble in water
- conduct electricity when liquid or in solution but do not when solid.

### Simple molecular (covalent)

Where found: in most non-metal elements and compounds.

Bonding: covalent bonds formed when atoms share pairs of electrons.

Structure: small, distinct groups of atoms.

Properties:

- low melting/boiling points
- a few are soluble in water
- most do not conduct electricity.

### Giant covalent

Where found: in a few non-metal elements and some compounds of non-metals.

Bonding: covalent bonds formed when atoms share pairs of electrons.

Structure: billions of atoms held together in a lattice structure.

Properties:

- high melting/boiling points
- insoluble in water
- most do not conduct electricity (except in carbon as graphite).

### Metallic

Where found: in all metals.

Bonding: metallic bonds are the electrostatic attraction between positive metal ions and negative delocalised electrons.

Structure: billions of ions held together in a giant lattice structure of positive ions in a 'sea' of negative delocalised electrons.

Properties:

- high melting/boiling points
- insoluble in water
- conduct electricity when solid or liquid.

The models help to explain some of the properties. For example:

- Substances with high melting points have many strong bonds that need to be broken during melting. Substances with low melting points have only weak forces that need to be overcome between molecules.
- Substances that conduct electricity have charged particles that can freely move. Substances that do not conduct electricity have charged particles that cannot move or particles that are not charged.



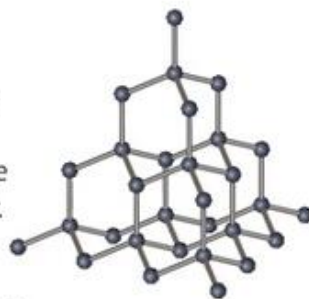
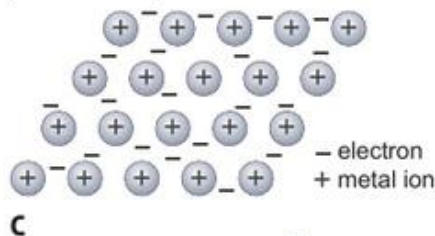


- 3** Explain why sodium metal conducts electricity when it is solid or liquid but sodium chloride conducts only when liquid.
- 4** A white solid with a melting point of  $2614^{\circ}\text{C}$  does not dissolve in water.
- a** Is the substance ionic or covalent or can you not tell? Explain your answer.
- b** Describe a test you could do to help you answer part a.
- 5** What do sodium ions and chloride ions have in common?

### Problems with bonding models

The bonding models that we use to explain the properties of matter all have certain weaknesses or limitations.

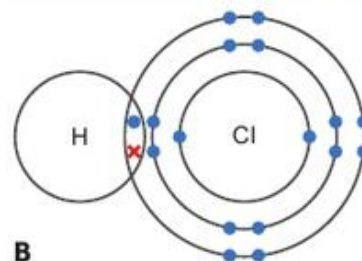
- The dot and cross diagram for hydrogen chloride in diagram B shows how electrons are shared in covalent bonds. However, dot and cross diagrams do not show the structure formed and they suggest that the electrons in different atoms are different, when they are actually all the same.
- The metallic model in diagram C shows the metal ions held in a lattice and explains why it conducts electricity, but the model does not show that the ions will be vibrating all the time.
- 3D ball and stick models, like diagram D for diamond, show which atoms are joined together and show the shape of the structure. However, they also show the atoms too far apart and there are not really 'sticks' holding the atoms together.



- 6 a** Draw a 3D structural ball and stick model of a sodium chloride lattice.
- b** Draw a dot and cross diagram for sodium chloride.
- c** Explain one strength and one weakness of a dot and cross diagram for sodium chloride.
- 7** Describe one strength and one weakness of the 3D models shown in photo A.

### Did you know?

Ideas about chemical bonds can be traced back to Roman times. The Roman poet Lucretius (c.95 BCE–c.55 BCE) imagined atoms as tiny spheres with fishhooks embedded in them. The atoms formed bonds when the fishhooks got tangled up with one another.



**B**

### Checkpoint

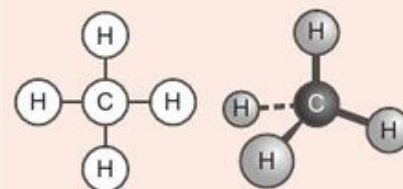
How confidently can you answer the Progression questions?

### Strengthen

- S1** Draw up a summary table describing the four 'types of bonding and structure' under the headings: 'where it occurs'; 'how bonds form'; 'type of structure'; 'properties' and 'examples'.

### Extend

- E1** The models below show methane.
- a** Draw a dot and cross diagram for methane.
- b** Describe one strength and one weakness of your dot and cross diagram and the two models below.



**E**

### Exam-style question

Suggest an explanation why the melting point of silicon dioxide ( $1610^{\circ}\text{C}$ ) is so much higher than the temperature at which solid carbon dioxide changes state ( $-78.5^{\circ}\text{C}$ ), although both compounds contain covalent bonds. (3 marks)

## Bonding in compounds

Sodium chloride and hydrogen chloride are both compounds of chlorine. Sodium chloride has a boiling point of  $1413^{\circ}\text{C}$ . Hydrogen chloride has a boiling point of  $-85^{\circ}\text{C}$ .

Explain this difference in boiling points in terms of the structure and bonding in the particles of the two compounds.

(6 marks)

### Student answer

As it is a metal and a non-metal, sodium chloride contains ionic bonds with lots of oppositely charged ions attracting each other to form a giant lattice structure [1]. Ionic compounds contain lots of strong bonds which keep it solid [2].



[3]

[1] This is a good description of the bonding and structure of sodium chloride.

[2] The question asked for an explanation of the high boiling point, not why it was solid.

[3] The diagram will get no credit as it is not labelled or described.

Hydrogen chloride contains covalent bonds and is made up of molecules. The covalent bonds are strong bonds that hold the atoms together in the molecules. Substances which contain molecules have low boiling points [4].

[4] This part of the answer does not explain why hydrogen chloride has a low boiling point.

### Verdict

This is an acceptable answer. It contains a good description of the bonding and structures of sodium chloride and hydrogen chloride. The answer has included information on both of the compounds in the question and uses correct scientific terminology.

This answer could be improved by explaining why the boiling points are low or high. The student should use scientific knowledge to explain the link between the structures of the compounds, the bonding within them and their boiling points.

#### Exam tip

You are more likely to get better marks in a question which requires extended writing if you plan your answer before you start. The plan can simply be a list of the keywords or phrases that you want to include. It is also useful to organise these keywords in a logical order to help structure your answer.





## Paper 1

### SC8 Acids and Alkalis

In the prehistoric world, lightning storms and volcanoes produced large quantities of gases that acidified our seas and lakes. Then, as now, acid rain was an environmental problem that caused extensive harm to plants and animals. We still live in a watery world, with many important chemical reactions occurring in aqueous solutions, where both the reactants and products dissolve in water. All solutions in water are either acidic, alkaline or neutral, as are many household and industrial chemicals. In this topic you will explore the nature of acidic and alkaline solutions, and investigate their most important reactions, properties and uses.

#### The learning journey

Previously you will have learnt at KS3:

- about solubility, solutes, solvents and solutions
- how common international hazard symbols are used
- about common acids, alkalis and neutral solutions
- about the use of indicators to test the pH of solutions
- about what happens during simple neutralisation reactions.

In this unit you will learn:

- about the ions in acids and alkalis, and how their concentrations are linked to pH
- what happens in the reactions between acids and different types of bases
- how different indicators can be used in acid-alkali titrations
- how different soluble and insoluble salts can be prepared in the laboratory.



# SC8a Acids, alkalis and indicators

Specification reference: C0.5; C3.1; C3.2; C3.3; **H** C3.4

## Progression questions

- Why are hazard symbols useful?
- What are the effects of acids and alkalis on some common indicators?
- What does the pH tell us about the ions in a solution?



**A** International hazard symbols are used in all countries.

All **aqueous solutions**, including those found in many household chemicals, are either **acidic**, **alkaline** or **neutral**. Some, including those that are neutral, can cause problems. For example, some acids and alkalis can be corrosive, toxic or harmful to the environment. The hazards associated with handling particular solutions are identified by international symbols, and they indicate the precautions that need to be taken when handling them.

- 1** Name two acids and two alkalis used in the home.
- 2**
  - a** Suggest an advantage of having internationally agreed hazard symbols.
  - b** Suggest a safety precaution for each of the hazard symbols shown in photo A.

## Indicators and pH

The **acidity** or **alkalinity** of a solution is measured on the **pH scale**. Most solutions lie between 0 and 14 on the scale. Solutions with a pH of 7 are neutral, while **acids** have a pH lower than 7 and **alkalis** have a pH greater than 7. The lower the pH, the more acidic the solution. The higher the pH, the more alkaline the solution.

The pH of a solution can be found by using **indicators** – substances that change colour depending on the pH. The **universal indicator** in photo B is made from a mixture of different indicators and produces a range of colours depending on the pH. Some other common indicators are shown in diagram C.



**B** Universal indicator can be used to find an approximate pH value.

- 3** Place the following pH values in order of increasing acidity. 4, 1, 7, 9, 6.
- 4**
  - a** What pH is the solution in photo B?
  - b** What colour would this solution turn each of the indicators in table C?

indicator	litmus	methyl orange	phenolphthalein
colour in alkaline solutions	blue	yellow	pink
colour in acidic solutions	red	red	colourless

**C** the colours of litmus, methyl orange and phenolphthalein

## Did you know?

Plant dyes can be used as pH indicators. The juice from cherries is bright red in acid solutions but turns blue/purple in alkalis. Curry powder contains the pigment curcumin, which changes from yellow at pH 7.4 to red at pH 8.6.

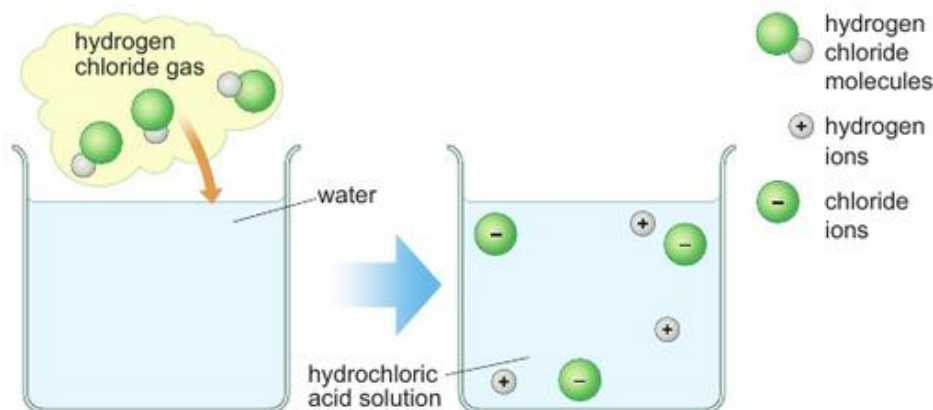


## Ions in acids and alkalis

An ion is an atom that has become charged by losing or gaining electrons. Losing electrons forms positive ions (e.g.  $\text{Na}^+$ ,  $\text{H}^+$ ). Gaining electrons forms negative ions (e.g.  $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{S}^{2-}$ ).

**Polyatomic ions** are formed when small groups of atoms, held together by covalent bonds, lose or gain electrons. Examples include  $\text{OH}^-$ ,  $\text{NO}_3^-$  and  $\text{SO}_4^{2-}$ .

Acids produce an excess of hydrogen ions ( $\text{H}^+$ ) when they dissolve in water. For example, hydrochloric acid is formed when hydrogen chloride gas dissolves in water and splits into  $\text{H}^+$  ions and  $\text{Cl}^-$  ions as shown in diagram E.



**E** Acids form hydrogen ions in water.

Alkalis produce excess hydroxide ions ( $\text{OH}^-$ ) in water. For example, solid sodium hydroxide splits into  $\text{Na}^+$  and  $\text{OH}^-$  ions when it dissolves.

### H

The higher the number of hydrogen ions in a certain volume, the higher their **concentration**. The higher their concentration, the more acidic the solution and the lower the pH.

The higher the concentration of hydroxide ions, the more alkaline the solution and the higher the pH.

Neutral solutions, such as pure water, have a pH of 7 and contain low, equal, concentrations of hydrogen ions and hydroxide ions.



- 8** Describe what happens to the acidity and pH of a solution, as more hydrogen ions are added.

### Exam-style question

- Explain why the pH of a hydrochloric acid increases when pure water is added. (2 marks)
- What would happen to the pH of salt solution (pH 7) if pure water was added? Briefly explain your answer. (2 marks)

Common acids	Formula
hydrochloric acid	$\text{HCl}$
sulfuric acid	$\text{H}_2\text{SO}_4$
nitric acid	$\text{HNO}_3$
Common alkalis	Formula
sodium hydroxide	$\text{NaOH}$
potassium hydroxide	$\text{KOH}$
calcium hydroxide	$\text{Ca(OH)}_2$

**D** common laboratory acids and alkalis



- 5** Describe what the pH scale tells us about acidity and alkalinity.

- 6** An acid is formed when hydrogen bromide dissolves in water.



- a** Suggest a possible pH for the solution and explain your choice.



- b** Write the symbols for the ions that form in this acidic solution.



- 7** What ions will each of the substances in table D produce when they dissolve in water?

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Describe the differences between an acidic and an alkaline solution.

### Extend

- E1** Explain how the nature of a solution is changing if the pH is: **a** increasing **b** decreasing.

# SC8b Looking at acids

Specification reference: **H** C3.5; **H** C3.7; **H** C3.8

## Progression questions

- H** What is the difference between dilute and concentrated solutions?
- H** How do changes in the concentration of hydrogen ions affect the pH of a solution?
- H** What is the difference between strong and weak acids?

**H**

- 1** 25 g of sulfuric acid is dissolved in water to make 200 cm<sup>3</sup> of concentrated sulfuric acid solution.



**a** What is a concentrated solution?



**b** What is the concentration of this solution in g dm<sup>-3</sup>?

A **concentrated** solution contains a lot of dissolved solute per unit volume, while a **dilute** solution contains only a small amount of solute.

## Worked example

$$\text{Concentration} = \frac{\text{amount dissolved}}{\text{volume of solution}}$$

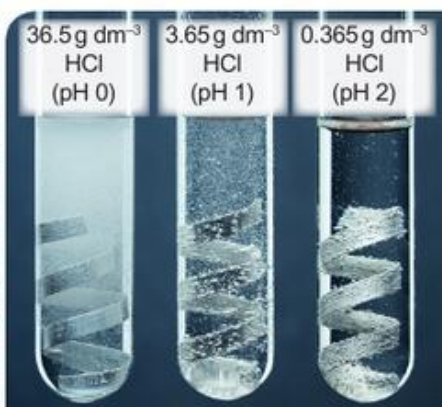
Units: grams per decimetre cubed, g dm<sup>-3</sup>

For example, if 4 g is dissolved in 50 cm<sup>3</sup>

$$\text{Concentration} = \frac{4}{0.05} = 80 \text{ g dm}^{-3}$$

Note: the minus sign shows that g is divided by dm<sup>3</sup>.  
1 dm<sup>3</sup> is the same volume as 1 litre or 1000 cm<sup>3</sup>.

Note: divide cm<sup>3</sup> by 1000 to change into dm<sup>3</sup>.



**A** More concentrated acids contain more hydrogen ions, so they react faster with magnesium metal.



**C** Universal indicator only gives a rough estimate of acidity. A **pH meter** is used for accurate pH measurements.

Accurate measurements of the pH of different concentrations of acid show that the concentration of hydrogen ions in an acid is linked to the pH. Table B shows that if the concentration of hydrogen ions is increased by a factor of 10, the pH decreases by 1. If the concentration decreases by a factor of 10, the pH increases by 1.

pH	0	1	2	3	4	5	6	7
difference in concentration of H <sup>+</sup> ions								

## B pH and concentration

So, hydrochloric acid with a pH of 0 is  $10 \times 10 \times 10 \times 10 = 10000$  times more acidic than vinegar, with a pH of 4. This means that a solution with a pH of 0 has a concentration of H<sup>+</sup> ions that is 10000 times greater than a solution with a pH of 4.



**2** What does the pH of a solution measure?



**3** An acid with a pH of 3 is diluted by a factor of 10. What will the new pH be?



**4** The most acidic rain recorded in Britain had a pH of 2. How much more acidic is this than normal rainwater with a pH of 5?

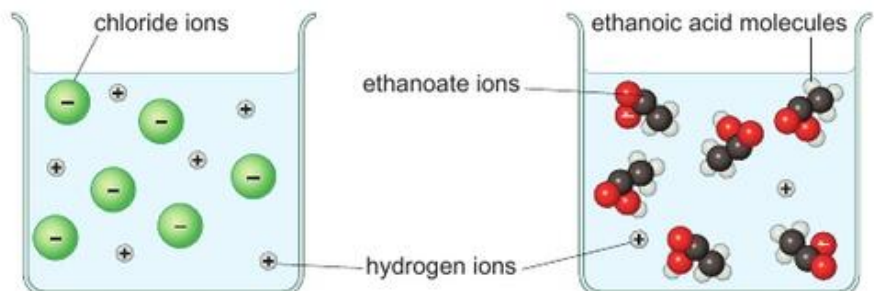


**5** What is the pH of the solution formed when 10 cm<sup>3</sup> of hydrochloric acid (pH 1) is made up to 1000 cm<sup>3</sup> with distilled water?



## H

The pH of an acid depends on the type of acid as well as its concentration. The acids in table D with low pH values are **strong acids**. Their molecules **dissociate** (break up) completely into ions when they dissolve in water and produce high concentrations of hydrogen ions. The other acids in the table are **weak acids**. They do not dissociate completely into ions in solution. Diagram E shows the difference between a strong and a weak acid.



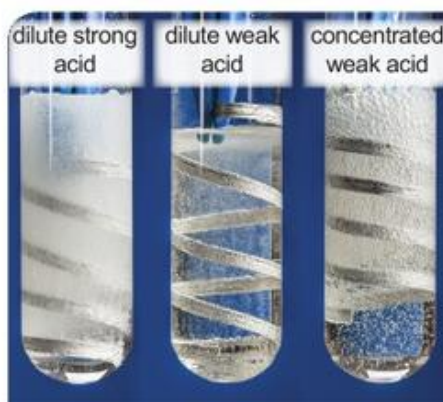
A strong acid like hydrochloric acid contains lots of hydrogen ions.

A weak acid like ethanoic acid contains only a few hydrogen ions.

## E strong and weak acids

The chemical properties of an acid depend on both the type of acid and its concentration, as shown in photo F.

- 6 List the strong and the weak acids in table D.
  - 7 About 1 in 60 000 butanoic acid molecules release hydrogen ions when they dissolve, while nearly all hydrogen iodide molecules do so.
- a Explain which acid is a weak acid.
  - b Explain which acid will have the higher pH, if both are the same concentration.



**F** A dilute solution of a strong acid can have a similar pH and reactivity as a concentrated solution of a weak acid. This is because they can have similar concentrations of hydrogen ions.

## Exam-style question

The pH of five solutions (J–N) are described below.

J = pH 4    K = pH 12    L = pH 7    M = pH 3    N = pH 1

- a Which two solutions have concentrations of hydrogen ions that differ by a factor of 10? (1 mark)
- b State and explain which solution could be a concentrated solution of a strong acid. (3 marks)

Acids (equal concentrations)	pH
carbonic acid	3.8
nitric acid	1.0
sulfuric acid	1.2
ethanoic acid	2.9
boric acid	5.2
hydrochloric acid	1.0

**D** the pH of acids of the same concentration

## Did you know?

The pH scale was developed in 1909 by Søren Sørensen (1868–1939) who was the head chemist in the Carlsberg Brewery in Copenhagen. The letters pH stand for 'power of hydrogen' where 'p' is short for the German word for power, potenz, and H is the symbol for hydrogen.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** What do the following terms tell you when they are used to describe an acid solution?
- a concentrated    b dilute  
c strong    d weak

## Extend

- E1** Explain how a concentrated solution of a weak acid could have the same pH and similar reactions to a dilute solution of a strong acid.

# SC8c Bases and salts

Specification reference: C3.9; C3.11; C3.13; C3.15

## Progression questions

- Why are metal oxides bases?
- What happens during neutralisation?
- How can a soluble salt be prepared from an acid and an insoluble base?



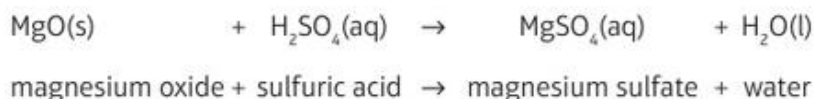
**A** Desulfurisation units attached to power station chimneys remove acidic sulfur-containing gases, by reacting them with the base calcium oxide.

- 1 What is a base?
- 2 Write a word equation and a symbol equation for the reaction of solid zinc oxide (ZnO) with sulfuric acid.

Acid	Salt formed
hydrochloric acid	chloride
sulfuric acid	sulfate
nitric acid	nitrate

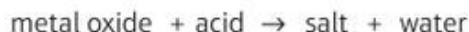
**C** salts from acids

**Bases** are substances that **neutralise** acids to form a **salt** and water only. All metal oxides are bases. For example, magnesium oxide is used to neutralise acids, like sulfuric acid, in industrial waste water.



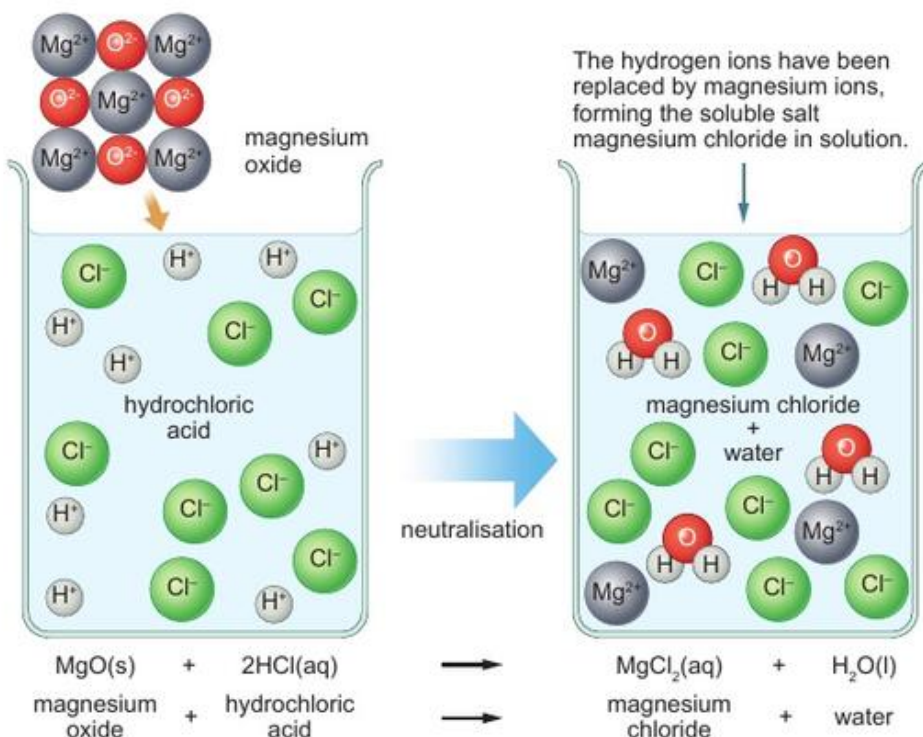
Note the use of **state symbols** in brackets after the formulae in symbol equations: (s) – solid; (l) – liquid; (g) – gas; (aq) – dissolved in water.

All neutralisation reactions with metal oxides occur in a similar way:



During neutralisation, hydrogen ions in the acid combine with oxide ions to form water. This removes the hydrogen ions and so the pH increases (becomes more neutral).

The salts are produced by replacing the hydrogen ions with metal ions, as shown in diagram B. Different acids form different salts, as shown in table C.



**B** how the salt magnesium chloride is formed





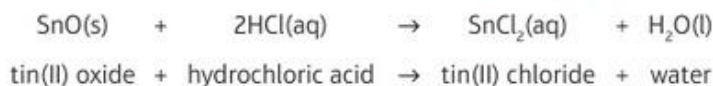
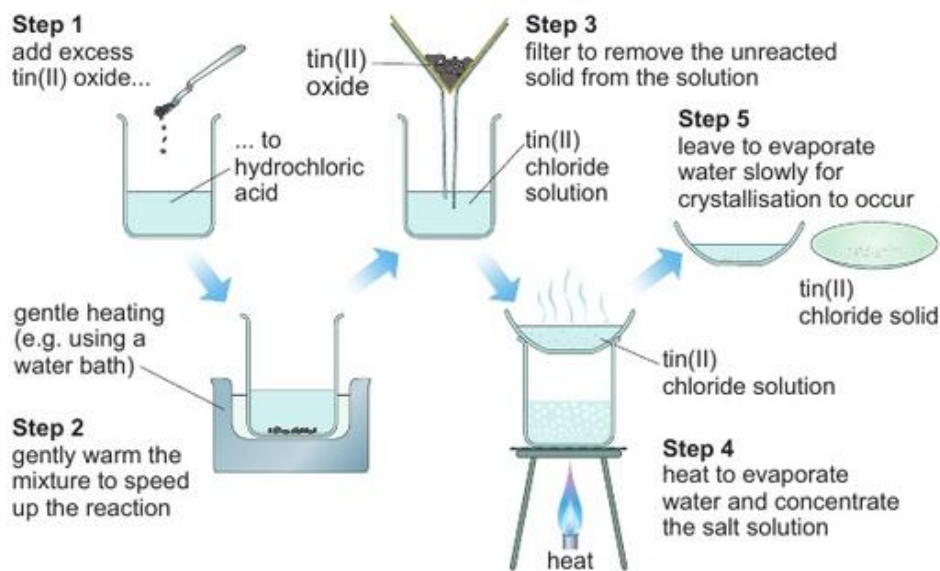
- 3** Write a word equation for the reaction between nitric acid and sodium oxide.
- 4** Copy and complete the symbol equation below, including the missing state symbols.



## Preparing soluble salts

The reaction between an acid and an insoluble metal oxide can be used to prepare samples of different soluble salts. For example, tin chloride can be prepared by reacting tin(II) oxide with hydrochloric acid solution. The steps involved are shown in diagram D.

An excess of the base is always added, to make sure that all the acid is used up. To make sure the prepared salt is pure, the mixture is **filtered** to remove the residue (the unreacted metal oxide) from the filtrate, leaving only the salt and water. A solid sample of the salt is then obtained by allowing the water to evaporate so that **crystallisation** of the salt occurs. Allowing the water to evaporate slowly will form larger crystals.



**D** the preparation of tin(II) chloride (eye protection must be worn)

- 5** Magnesium nitrate can be made by reacting magnesium oxide with an acid.

- a** Name the acid used.
- b** Why should the acid solution be warmed slightly?
- c** Why is the magnesium oxide added in excess?
- d** Why is the mixture filtered?
- e** Why is the salt solution heated?
- 6** Explain what happens to the hydrogen ions during the reaction between hydrochloric acid and tin(II) oxide.

## Did you know?

'Limelight', meaning 'the centre of attention', originated from 19th-century stage lighting systems. These used burners to heat lumps of lime (calcium oxide), making them glow white hot.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** Draw a flowchart to explain the steps involved in preparing a soluble salt from an insoluble base.

## Extend

**E1** Aluminium nitrate,  $\text{Al}(\text{NO}_3)_3$ , can be made using aluminium oxide,  $\text{Al}_2\text{O}_3$ , and nitric acid,  $\text{HNO}_3$ . Describe the preparation of aluminium nitrate and explain each step of the process, including word and symbol equations.

## Exam-style question

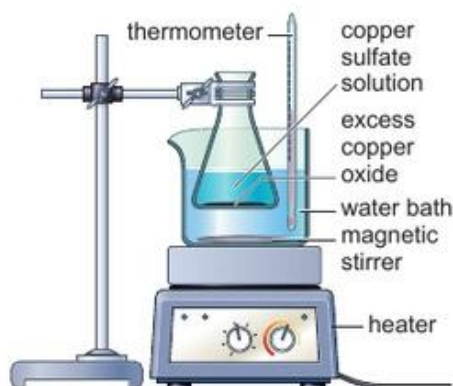
Explain, with reference to the ions present, how the pH of the acidic chimney gases in photo A will change as they pass over calcium oxide. (3 marks)

# SC8c Core practical – Preparing copper sulfate

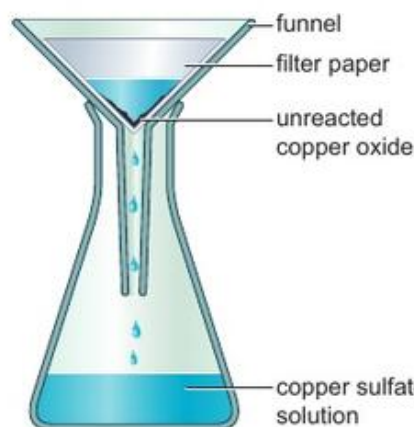
Specification reference: C3.17

## Aim

Investigate the preparation of pure, dry, hydrated copper sulfate crystals starting from copper oxide including the use of a water bath.



**A** Heating the acid makes the reaction between sulfuric acid and copper oxide faster.

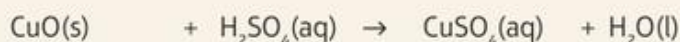


**B** Filtration separates the excess copper oxide (the residue) from the copper sulfate solution (the filtrate).

Salts, such as copper sulfate, are compounds formed by reacting an acid with a base. In the formation of a salt, the hydrogen ions in the acid are replaced by a positive ion (usually a metal ion) from the base. Different acids and bases produce different salts with a variety of properties and uses. For example, iron(III) oxide and hydrochloric acid form the salt iron(III) chloride ( $\text{FeCl}_3$ ), which is used in water treatment. Copper sulfate ( $\text{CuSO}_4$ ) is used by farmers and gardeners as a fungicide and herbicide. Skilled chemists can prepare, purify and obtain samples of many different salts.

## Your task

Copper oxide reacts with warm sulfuric acid to produce a blue solution of the salt copper sulfate. You are going to use these reactants to prepare pure, dry hydrated copper sulfate crystals.



copper oxide + sulfuric acid  $\rightarrow$  copper sulfate + water

Copper oxide (the base) is insoluble in water. So when all the sulfuric acid has been used up, the excess copper oxide remains. Excess copper oxide is added to make sure that all the acid is used up. The remaining copper oxide is then separated from the copper sulfate solution by filtration. If the water is allowed to evaporate, a pure, dry sample of copper sulfate can be obtained.

## Method

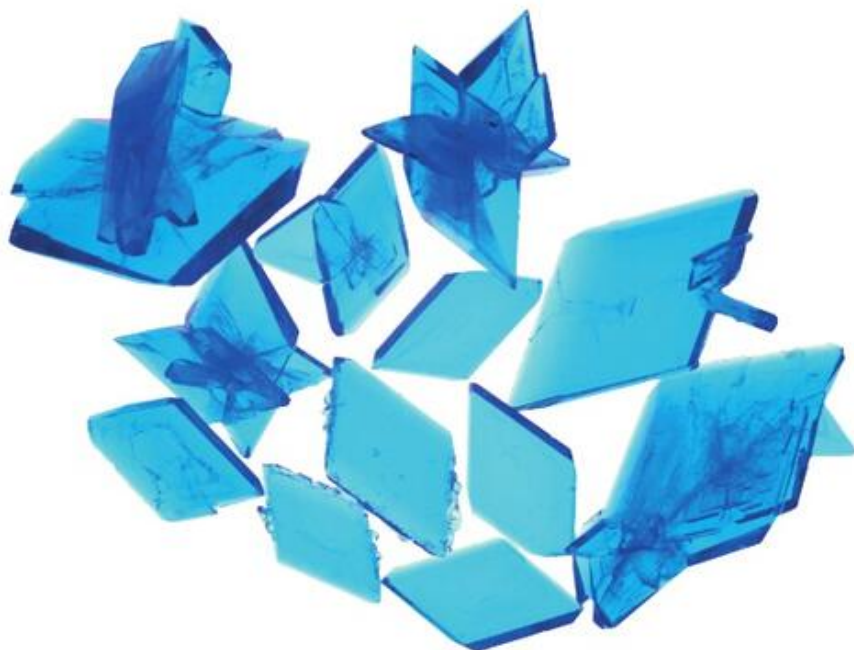
Wear eye protection.

- Measure  $20\text{ cm}^3$  of dilute sulfuric acid using a measuring cylinder and pour it into a small conical flask.
- Warm the acid in a water bath set at  $50^\circ\text{C}$ . Use a thermometer to measure the temperature.
- Add a little copper oxide powder to the acid and stir.
- If all the copper oxide reacts, and disappears, add a little more. Stop when the copper oxide is in excess and no longer reacts.
- Filter the mixture and transfer the filtrate to an evaporating basin.
- Heat the evaporating basin by placing it over a beaker of water heated with a Bunsen burner as shown in diagram D on the previous page. Stop heating when crystals start to form.
- Pour the solution into a watch glass and leave for a few days to allow all the water to evaporate.



## Exam-style questions

- 1 State why copper sulfate is described as a salt. (1 mark)
- 2 State why the sulfuric acid is heated in step B. (1 mark)
- 3 In step E, explain why the copper oxide gets stuck in the filter paper while the copper sulfate goes through it. (2 marks)
- 4 Explain why a water bath is used in step B rather than heating with a Bunsen burner. (2 marks)
- 5 Explain how you know a chemical reaction has occurred in step C. (2 marks)
- 6 State how you know when all the acid has been used up. (1 mark)
- 7 Give a reason why it is important to make sure the copper oxide is in excess in step D. (1 mark)
- 8 Describe one safety precaution that should be taken during this experiment and explain why it is necessary. (2 marks)
- 9 Nickel chloride ( $\text{NiCl}_2$ ) is a soluble salt. It can be made by reacting insoluble nickel oxide ( $\text{NiO}$ ) with hydrochloric acid ( $\text{HCl}$ ).
  - a Write a word equation for this reaction. (1 mark)
  - b Write a balanced equation with state symbols. (2 marks)
  - c Briefly describe the three main stages involved in preparing a pure, solid sample of nickel chloride. (3 marks)
- 10 Two class groups prepared some zinc chloride. One group produced lots of very small crystals while the other group produced larger crystals. Suggest an explanation for the groups producing different-sized crystals. (2 marks)



**C** Large crystals of copper sulfate form if the water evaporates slowly from a concentrated solution of copper sulfate.

# SC8d Alkalis and balancing equations

Specification reference: C0.1; C0.2; C0.3; C3.10; C3.11

## Progression questions

- What are alkalis?
- What happens when alkalis react with acids?
- How do we balance chemical equations?



**A** These skin burns were caused by kneeling in wet cement, which contains an alkali called calcium hydroxide.

A base is any substance that reacts with an acid to form a salt and water only. Many bases are insoluble in water – they do not dissolve. A base that *can* dissolve in water, a *soluble* base, is called an **alkali**. Alkalis form alkaline solutions with pH values above 7.



- 1** Explain why all alkalis are bases, but not all bases are alkalis.

## Common alkalis

Copper(II) hydroxide and most other metal hydroxides are insoluble bases.

## Did you know?

Magnesium hydroxide is sparingly soluble. Just enough magnesium hydroxide dissolves to make an alkaline solution. It forms a cloudy white suspension in water, which is used as a medicine to neutralise excess stomach acid.



**B**

However, these bases are soluble and therefore also alkalis:

- sodium hydroxide, NaOH, and other group 1 hydroxides
- calcium hydroxide, Ca(OH)<sub>2</sub>, and other group 2 hydroxides.

Notice that the chemical formulae for these alkalis look different. In general they are MOH for group 1 hydroxides, where M stands for the metal's chemical symbol, but M(OH)<sub>2</sub> for group 2 hydroxides. This is because the ions formed by group 2 atoms have a 2+ charge, but hydroxide ions have a 1– charge. The brackets show that two OH<sup>–</sup> ions are needed to produce a neutral compound.

- 2** Write the formula for:



- a** lithium hydroxide      **b** magnesium hydroxide.

## Modelling reactions of alkalis with acids

Like other bases, alkalis react with acids to produce salts and water only. For example:



During the neutralisation reaction, the reaction mixture becomes warmer. When an alkali is added to an acid, the pH increases and may go higher than 7 if enough alkali is added.



You can model the reaction using a **balanced equation**. In such equations, the numbers of atoms of each element must be the same on both sides of the arrow.

NaOH(aq)	+	HCl(aq)	→	NaCl(aq)	+	H <sub>2</sub> O(l)
1				Na	1	
1				O		1
1		1		H		2
		1		Cl	1	

**C** There is one Na, one O, two H and one Cl atom on each side of this equation.

The water is formed from OH in the alkali and H in the acid.

You may need to write a number in front of one or more formulae to balance an equation. Remember that the formulae themselves cannot be changed. For example:

sodium hydroxide + sulfuric acid → sodium sulfate + water

2NaOH(aq)	+	H <sub>2</sub> SO <sub>4</sub> (aq)	→	Na <sub>2</sub> SO <sub>4</sub> (aq)	+	2H <sub>2</sub> O(l)
(2 × 1) = 2				Na	2	
(2 × 1) = 2		4		O	4	(2 × 1) = 2
(2 × 1) = 2		2		H		(2 × 2) = 4
		1		S	1	

**D** The number 2 in front of NaOH and H<sub>2</sub>O in the balanced equation shows that two units of these substances are needed.

Equations for the reactions of group 2 hydroxides with acids may look more complex. For example:

calcium hydroxide + nitric acid → calcium nitrate + water

Ca(OH) <sub>2</sub> (aq)	+	2HNO <sub>3</sub> (aq)	→	Ca(NO <sub>3</sub> ) <sub>2</sub> (aq)	+	2H <sub>2</sub> O(l)
1				Ca	1	
(1 × 2) = 2		(2 × 3) = 6		O	(3 × 2) = 6	(2 × 1) = 2
(1 × 2) = 2		(2 × 1) = 2		H		(2 × 2) = 4
		(2 × 1) = 2		N	(1 × 2) = 2	

**E** The number 2 outside the brackets for OH and NO<sub>3</sub> shows that there are two of these groups of atoms in the formulae for these substances.

- 6** Balance this equation for the reaction between potassium hydroxide solution and phosphoric acid:
- $$\text{KOH(aq)} + \text{H}_3\text{PO}_4\text{(aq)} \rightarrow \text{K}_3\text{PO}_4\text{(aq)} + \text{H}_2\text{O(l)}$$

- 3** Potassium hydroxide, KOH, reacts with hydrochloric acid in a similar way to sodium hydroxide. Write the balanced equation for this reaction, including state symbols.

- 4** Explain why the equation in diagram D is *balanced*.

- 5** Balance this equation for the reaction between barium hydroxide solution and sulfuric acid:
- $$\text{Ba(OH)}_2\text{(aq)} + \text{H}_2\text{SO}_4\text{(aq)} \rightarrow \text{BaSO}_4\text{(s)} + \text{H}_2\text{O(l)}$$

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** What happens when metal hydroxides react with acids?  
**S2** How are chemical equations balanced?

### Extend

- E1** Evaluate the use of balanced equations to model the reactions of metal hydroxides with acids.

### Exam-style question

Magnesium hydroxide solution reacts with stomach acid, which is hydrochloric acid. Magnesium chloride solution is a product. Write a balanced equation for the reaction, including state symbols. (3 marks)

# SC8d Core practical – Investigating neutralisation

Specification reference: C3.6

## Aim

Investigate the change in pH on adding powdered calcium hydroxide or calcium oxide to a fixed volume of dilute hydrochloric acid.



**A** Indigestion liquids and tablets contain different antacids.

Stomach acid contains hydrochloric acid. Acid indigestion causes a burning feeling in the chest and throat. Antacids are substances that neutralise stomach acid to relieve indigestion. Magnesium hydroxide is a white solid used in antacid tablets. It is sparingly soluble in water, so it is also used as a white suspension that some people find easier to swallow.

Magnesium hydroxide neutralises hydrochloric acid:

magnesium hydroxide + hydrochloric acid → magnesium chloride + water



## Your task

Calcium and magnesium are in group 2 of the periodic table. Calcium hydroxide has similar chemical properties to magnesium hydroxide, but it is more soluble in water. You will investigate the change in pH when you add powdered calcium hydroxide to dilute hydrochloric acid. You will add small portions of powder to the acid and record the pH of the mixture after each addition.

## Method

Wear eye protection.

- A** Use a measuring cylinder to add 50 cm<sup>3</sup> of dilute hydrochloric acid to a beaker.
- B** Estimate and record the pH of the contents of the beaker.
  - Put a piece of universal indicator paper onto a white tile.
  - Dip the end of a glass rod into the liquid, then tap it onto the universal indicator paper.
  - Wait 30 seconds, then match the colour to the appropriate pH on a pH colour chart.
  - Rinse the glass rod with water.
- C** Measure out 0.3 g of calcium hydroxide powder onto a piece of paper or a 'weighing boat'.
- D** Add the calcium hydroxide powder to the beaker and stir. Then estimate and record the pH of the mixture.
- E** Repeat steps B and C seven times so that you add a total of 2.4 g of calcium hydroxide powder to the acid.
- F** Plot a graph with pH on the vertical axis and mass of calcium hydroxide on the horizontal axis.

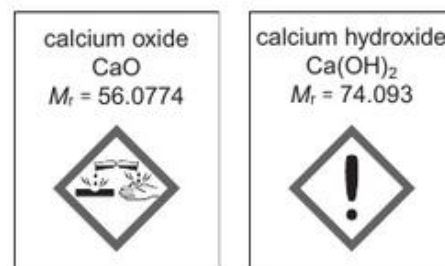


**B** A pH meter may also be used to measure pH.



## Exam-style questions

- 1
  - a Name the soluble salt formed when hydrochloric acid reacts with calcium hydroxide. (1 mark)
  - b Write the balanced equation, including state symbols, for the reaction between calcium hydroxide powder and dilute hydrochloric acid. (3 marks)
- 2 Diagrams C and D show part of the labels found on two laboratory containers.
  - a Give two reasons why hazard symbols are used on chemical containers. (2 marks)
  - b Explain why it may be more hazardous to handle calcium oxide than calcium hydroxide. (3 marks)
- 3 Give two reasons that explain why eye protection must be worn when using dilute hydrochloric acid. (2 marks)
- 4 A student investigates the change in pH when calcium hydroxide powder is added to 100 cm<sup>3</sup> of dilute hydrochloric acid.
  - a State two control variables in his experiment. (2 marks)
  - b State the independent variable in his experiment. (1 mark)
  - c Describe how the student could modify his experiment to investigate temperature changes instead of pH changes. (1 mark)
- 5 A student wants to find the mass of calcium oxide powder that produces a neutral solution when added to 75 cm<sup>3</sup> of dilute hydrochloric acid. She adds 0.5 g portions of the powder to the acid, and measures the pH each time with a pH meter. Table E shows her results.
  - a Predict the mass of calcium oxide that produces a neutral solution. (1 mark)
  - b State and explain how the student could improve her experiment to obtain a more accurate result. (3 marks)
- 6 The pH of a solution may be determined using universal indicator paper or using a pH meter.
  - a State why a pH meter must be calibrated using a solution with a known pH value. (1 mark)
  - b Explain whether indicator paper or a pH meter has the higher resolution. (2 marks)



C

D

Mass of calcium oxide added (g)	pH of reaction mixture
0.0	0.3
0.5	0.6
1.0	2.8
1.5	12.3
2.0	12.5

E

# SC8e Alkalis and neutralisation

Specification reference: C3.11; C3.14; C3.16; C3.18

## Progression questions

- What happens to the ions from acids and alkalis during neutralisation?
- What is titration?
- How do we make a soluble salt using titration?



**A** Potassium sulfate, manufactured using sulfuric acid and a soluble reactant, is a soluble salt used in fertilisers.

Soluble bases or alkalis react with acids to form a salt and water only.  
The type of reaction involved is neutralisation.

## Did you know?

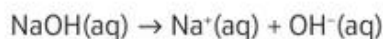
The Russian government imposed a high tax on table salt, sodium chloride, in the 17th century. This led to a 'salt riot' in Moscow in 1648.

## Ions and neutralisation

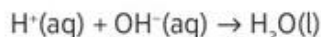
Hydrochloric acid and other acids are a source of hydrogen ions,  $\text{H}^+(\text{aq})$ , in solution:



Sodium hydroxide and other alkalis are a source of hydroxide ions,  $\text{OH}^-(\text{aq})$ , in solution:



In a neutralisation reaction, hydrogen ions from the acid react with hydroxide ions from the alkali. Water, a simple molecular substance containing covalent bonds, is formed in the reaction:



The other ions from the acid and alkali stay in the solution as ions of the dissolved salt. For example,  $\text{Na}^+(\text{aq})$  ions from sodium hydroxide and  $\text{Cl}^-(\text{aq})$  ions from hydrochloric acid remain after neutralisation. These ions combine to form solid sodium chloride,  $\text{NaCl}(\text{s})$ , when the water evaporates.

**6<sup>th</sup>** 1 Explain how water is formed in neutralisation reactions.

**7<sup>th</sup>** 2 Explain, in terms of ions, how potassium hydroxide solution reacts with sulfuric acid to form water and potassium sulfate.



## Soluble salts from alkalis

You can obtain a dry soluble salt from its solution by crystallisation. It is important to have a neutral solution before evaporating the water, otherwise you will contaminate the salt with an excess of one reactant.



- 3 Other than the hazards caused by hot liquids and solids, suggest a hazard caused by crystallising a salt from a solution that contains excess acid or alkali.

To obtain a neutral solution you need to mix an acid and an alkali in the correct proportions so that you end up with a solution that contains only water and the desired salt. You can do this using **titration**.

## Titration

In a titration, acid is added from a **burette** to a fixed volume of alkali in a conical flask. The burette is a tall piece of glassware with  $0.1 \text{ cm}^3$  graduations. You control the flow using a tap at the bottom. It is possible to add just one drop at a time.

You could use a measuring cylinder to measure out the alkali, but a **pipette** provides more accurate and repeatable measurements. A few drops of indicator are added to the alkali so you can follow the reaction. The **end-point** is when the indicator changes colour. Single indicators such as methyl orange or phenolphthalein are used because their obvious colour changes give you a sharp end-point.

To make a pure, dry salt:

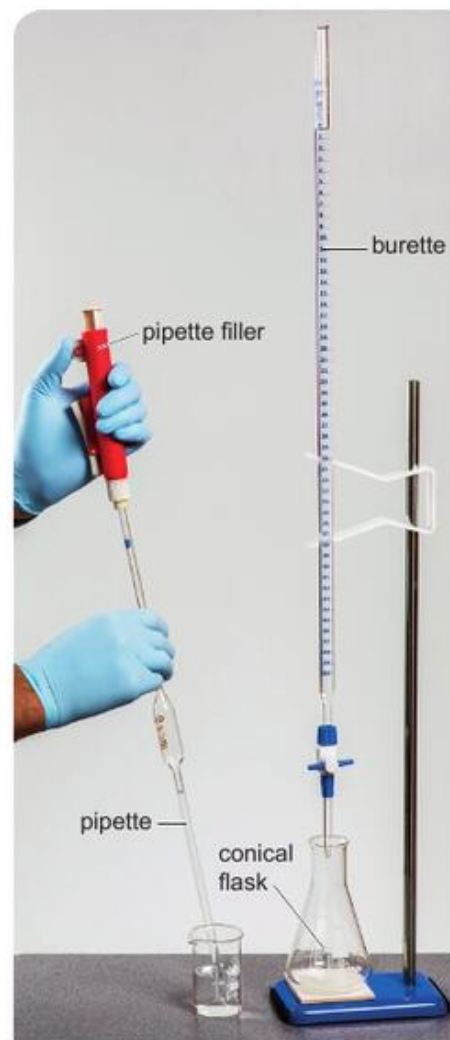
- carry out a titration
- note the exact volume of acid needed to neutralise the alkali
- use the burette to add the correct volume of acid *without* the indicator
- evaporate the water from the solution formed.



- 4 In a titration, name the most suitable apparatus to add alkali to the flask, and to add acid to the alkali.



**B** producing a dry salt by crystallisation using a hot water bath



**C** This apparatus is used to carry out a titration. A white tile underneath the conical flask makes the end-point easier to see.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** Describe how titration is used to prepare soluble salts.

## Extend

**E1** Evaluate the preparation of sodium and potassium salts using titration rather than by the reactions of the metals with acids.

## Exam-style question

- a** Give a reason why universal indicator is not a good choice of indicator to use in a titration. (1 mark)
- b** Suggest a suitable indicator to use instead. (1 mark)

# SC8f Reactions of acids with metals and carbonates

Specification reference: **H** C0.4; C3.11; C3.12

## Progression questions

- What happens when an acid reacts with a metal?
- What happens when an acid reacts with a metal carbonate?
- What are the tests for hydrogen and carbon dioxide?



**A** Hydrogen gas is less dense than air and so was used to fill early air ships. Unfortunately it is also very flammable.



**B** Hydrogen gas is colourless and odourless. When hydrogen is mixed with air, it is highly flammable. To test for hydrogen, place a lighted splint in a hydrogen and air mixture. A squeaky pop is heard.

- 2** Zinc reacts with dilute hydrochloric acid. Write:



**a** the word equation



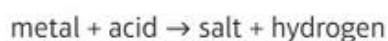
**b** the balanced equation



**H c** the half equation to show what happens to the zinc atoms.

## Acids and metals

Some metals, such as copper and silver, do not react with dilute acids. Metals such as potassium and sodium react explosively with dilute acids. Metals in the middle of the **reactivity series**, such as magnesium and zinc, react steadily with dilute acids. **Effervescence** is seen as hydrogen gas bubbles are produced. The reaction also produces a salt, giving the general reaction:



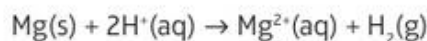
Magnesium reacts with sulfuric acid to form magnesium sulfate, and with hydrochloric acid to form magnesium chloride. The first name of the salt comes from the metal and the second name comes from the acid.



- 1** Explain why gold sulfate is not formed when gold is added to dilute sulfuric acid.

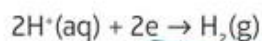
### H

All acids form hydrogen ions,  $\text{H}^+$ , in aqueous solution. The metal atoms react with the hydrogen ions to form metal ions and hydrogen molecules. This can be summarised in an **ionic equation**, in which we only show the ions that change in the reaction. For example:

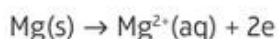


The other ions from the acid (e.g. sulfate, chloride) do not change during the reaction. They are known as **spectator ions**.

The hydrogen ions gain electrons to form hydrogen molecules. We can show what happens to electrons in a **half equation**:



The magnesium atoms lose electrons:



A loss of electrons is an **oxidation** reaction. A gain of electrons is a **reduction** reaction. You can remember this using the mnemonic OILRIG. (Oxidation Is Loss, Reduction Is Gain).

e represents an electron.  
It is not necessary to include the negative charge.

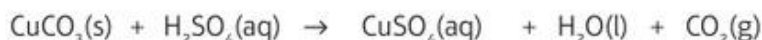


## Acids and carbonates

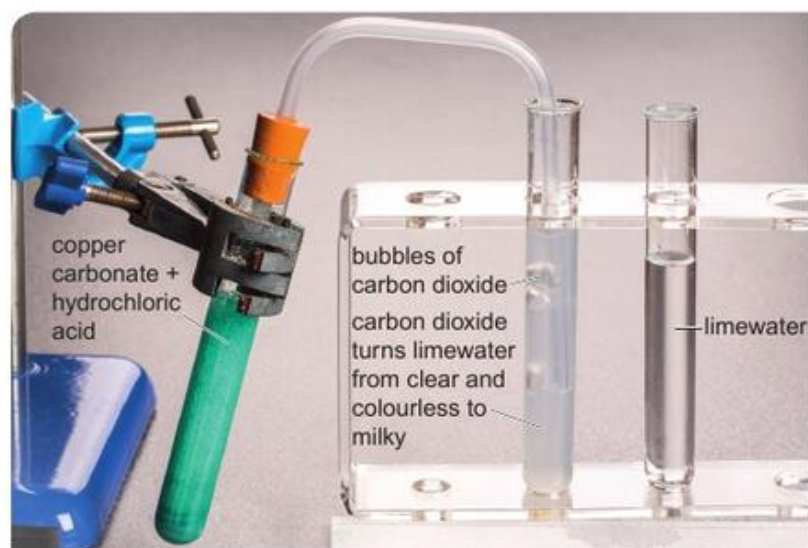
Acids react with metal carbonates to form a salt, water and carbon dioxide. Bubbles of carbon dioxide are produced and the solid metal carbonate disappears if there is enough acid to react with it.



Copper carbonate reacts with sulfuric acid to form copper sulfate, with hydrochloric acid to form copper chloride, and with nitric acid to form copper nitrate.



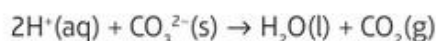
During these reactions, the hydrogen ions from the acids react with the carbonate ions to form water and carbon dioxide molecules.



**C** Carbon dioxide is another colourless and odourless gas. To test for carbon dioxide, bubble the gas through limewater (calcium hydroxide solution). The limewater turns milky if carbon dioxide is present.

**H**

The sulfate, chloride and nitrate ions are spectator ions. The ionic equation is:



**8<sup>th</sup>**

- 6** Write the ionic equation for the reaction between magnesium carbonate and hydrochloric acid.

### Did you know?

Carbon dioxide is denser than air and does not support burning. This makes it useful in fire extinguishers.

**8<sup>th</sup>**

- 3** Give the formula of the salt formed when solid calcium carbonate,  $\text{CaCO}_3$ , reacts with dilute hydrochloric acid.

**6<sup>th</sup>**

- 4** Explain the difference between the state symbols (aq) and (l).

**9<sup>th</sup>**

- 5** Write the balanced equation for the reaction between zinc carbonate,  $\text{ZnCO}_3$ , and nitric acid,  $\text{HNO}_3$ .

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Describe a reaction to make:  
**a** hydrogen  
**b** carbon dioxide.

Include an equation for each reaction and a description of how to test for each gas.

### Extend

- E1** **H** Dilute hydrochloric acid is added to zinc powder. Write word, balanced and ionic equations for the reaction that occurs. Include state symbols.

### Exam-style question

Describe a test to show that a gas is hydrogen.

(2 marks)

# SC8g Solubility

Specification reference: **H** C0.4; C3.19; C3.20; C3.21

## Progression questions

- What are the rules for solubility of common substances in water?
- How do you prepare a sample of a pure, dry insoluble salt?
- How do you predict whether a precipitate will be formed in a reaction?



**A** A red precipitate of silver chromate forms when sodium chromate solution is added to silver nitrate solution.

- 1** Which of the following substances are soluble in water and which are insoluble? Sodium chloride, lead nitrate, calcium sulfate, potassium hydroxide, silver chloride, calcium carbonate, ammonium carbonate.

- 2** Write a solubility rule for lead compounds.

## Did you know?

The reaction between sodium chromate and silver nitrate is used to stain nerve cells for viewing under a microscope.

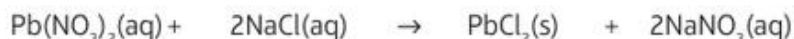
A **precipitation** reaction is one in which soluble substances in solutions cause an insoluble **precipitate** to form. Photo A shows an example. Table B shows some general rules that can be used to predict whether precipitates will form.

Soluble in water	Insoluble in water
all common sodium, potassium and ammonium salts	
all nitrates	
most chlorides	silver, lead chlorides
most sulfates	lead, barium, calcium sulfates
sodium, potassium and ammonium carbonates	most carbonates
sodium, potassium and ammonium hydroxides	most hydroxides

## B some solubility rules

Solutions of lead nitrate and sodium chloride react to form soluble sodium nitrate and a white precipitate of lead chloride. The state symbols in the balanced equation show which substances are soluble and which are insoluble.

lead nitrate + sodium chloride  $\rightarrow$  lead chloride + sodium nitrate



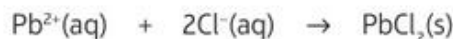
The test for carbon dioxide using limewater (see CC8f) is also a precipitation reaction, in which insoluble calcium carbonate is formed:

calcium hydroxide (limewater) + carbon dioxide  $\rightarrow$  calcium carbonate + water



## H

Ionic equations are also used to show the formation of precipitates. The formulae of the ions that react together to form the precipitate are shown on the left, and the formula of the precipitate is shown on the right. The ionic equation for the formation of lead chloride is:



The sodium and nitrate ions do not change, so they are spectator ions.



All salts are ionic. When two solutions containing soluble salts react together, the ions from the salts swap. For example:



We can predict whether a precipitate will form by checking the solubilities of the products. If both products are soluble, no precipitate will form.

In the word equation above, potassium sulfate does not form a precipitate because all potassium salts are soluble. However, copper carbonate is not in the list of soluble carbonates, so it should form as a precipitate:

**H**

The ionic equation is:  $\text{Cu}^{2+}(\text{aq}) + \text{CO}_3^{2-}(\text{aq}) \rightarrow \text{CuCO}_3(\text{s})$

## Preparation of insoluble salts

A pure, dry sample of an insoluble salt can be prepared from two soluble salts in this way:

- Wear eye protection.
- Mix the two solutions in a beaker, then filter the mixture.
- Rinse the beaker with a little distilled water and pour this through the funnel.
- Pour a little distilled water over the precipitate in the funnel.
- Carefully remove the filter paper containing the precipitate and dry it in a warm oven.

**5** Explain:



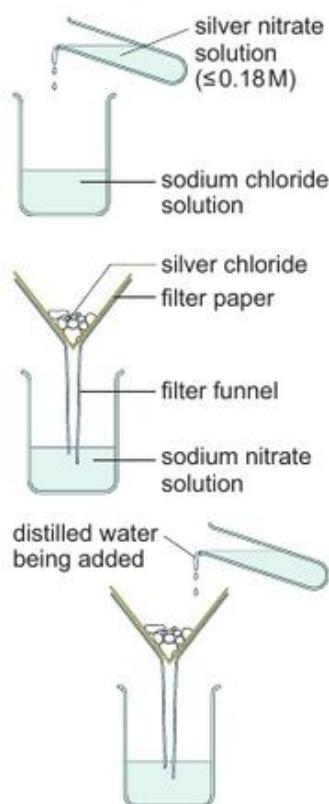
**a** why the beaker is rinsed with a little distilled water and the washings are poured through the funnel



**b** why distilled water is poured over the precipitate in the funnel.



**6** Suggest two other ways of drying the precipitate.



**C** preparing an insoluble salt

**3** Magnesium sulfate solution reacts with barium chloride solution.



**a** Predict the name of the precipitate, if any.



**b** Write the balanced equation. Include state symbols.



**H c** Write the ionic equation for this reaction. Include state symbols.

**4** Sodium chloride solution is added to copper nitrate solution.



**a** Predict the name of the precipitate, if any.



**b** Justify your answer.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** Describe how to prepare a pure, dry sample of lead sulfate from two named solutions. Write a balanced equation for the reaction. Include state symbols.

## Extend

**E1 H** The following aqueous solutions are available: sodium carbonate, copper sulfate, barium chloride. Any two of the solutions can be mixed together at a time. Predict which combinations of these solutions will produce a precipitate and name the precipitate. Write an ionic equation, including state symbols, for each reaction.

## Exam-style question

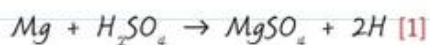
Zinc carbonate is an insoluble salt. Describe how you would use solutions of potassium carbonate and zinc chloride to produce a pure, dry sample of zinc carbonate. (4 marks)

## Magnesium sulfate

Soluble salts can be made by reacting an acid with an insoluble metal compound.

Plan an experiment to prepare pure, dry crystals of magnesium sulfate,  $\text{MgSO}_4$ , from a magnesium compound and a suitable acid. Start by choosing suitable reactants to use; you may wish to write a balanced equation to help you with your plan. (6 marks)

### Student answer



I will make magnesium sulfate from magnesium ribbon and sulfuric acid [2].

Place  $25 \text{ cm}^3$  of the acid in a beaker. Add magnesium ribbon a piece at a time until the bubbles of hydrogen stop. Filter the mixture and collect the solution in an evaporating basin [3]. Heat the basin until all the water has evaporated, leaving pure, dry crystals of magnesium sulfate [4].

[1] The balanced equation is incorrect. Hydrogen gas exists as  $\text{H}_2$  molecules not separate hydrogen atoms.

[2] Sulfuric acid is correct, but the question asks for a magnesium compound (such as magnesium oxide). Magnesium is an element.

[3] The answer could also have explained that you need to filter the mixture to remove any excess magnesium.

[4] The last step is incorrect. If you evaporate all of the water, you will be left with a dry powder.

### Verdict

This is an acceptable answer. It contains some correct chemistry, and the steps of the practical are written in a clear and logical order.

The answer could be improved by using a magnesium compound as mentioned in the question. The answer would also be better if it had a correct balanced equation, and described a suitable method for allowing crystals to form from the solution.

#### Exam tip

Make sure you revise the Core Practicals. The method asked for here is very similar to the preparation of copper sulfate which is a Core Practical.





## Paper 1 and Paper 2

### SC9 Calculations Involving Masses

In most chemical reactions it is important to mix the reactants in the correct amounts to form the maximum amount of product and to avoid waste. Iron is produced from iron ore in a blast furnace. Every tonne of iron ore needs to be mixed with 850 kg of limestone and 50 kg of coal. This should produce 350 kg of iron and 910 kg of a substance called 'slag' that can be used in concrete and for insulation. In this unit you will be learning how to calculate these amounts.

#### The learning journey

Previously you will have learnt at KS3:

- how to represent elements and compounds using symbols
- how mass is conserved during changes of state and chemical reactions
- how to show chemical reactions using equations.

In this unit you will learn:

- how to use relative atomic masses to calculate relative formula masses of elements and compounds
- how to work out empirical and molecular formulae of compounds
- how to calculate the mass of reactants or products in a reaction
- how to calculate the concentration of a solution
- **H** about the Avogadro constant and the quantity 1 mol of a substance
- **H** how to calculate the numbers of particles in a substance.

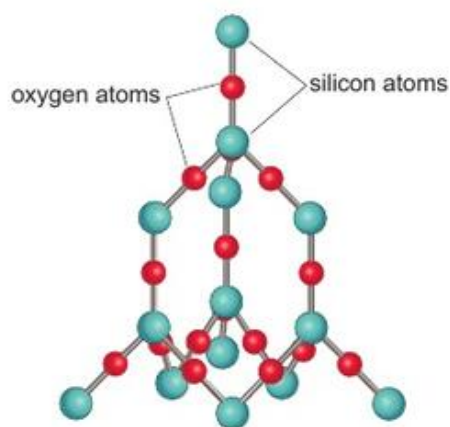


# SC9a Masses and empirical formulae

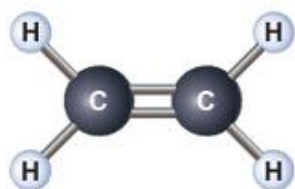
Specification reference: C1.43; C1.44; C1.45; C1.46

## Progression questions

- How do you calculate the relative formula mass of a compound?
- What is the difference between an empirical formula and a molecular formula?
- How do you determine the empirical formula of a compound?



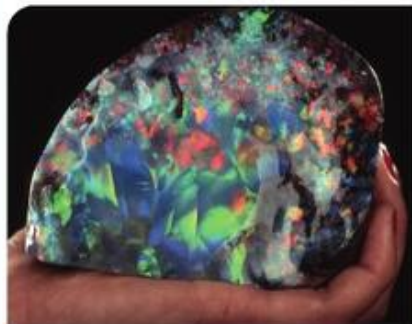
**A** Silicon dioxide is a giant, covalent lattice structure (of billions of atoms). Its empirical formula is  $\text{SiO}_2$  (the ratio of silicon atoms to oxygen atom is 1:2).



**B** Ethene has the molecular formula  $\text{C}_2\text{H}_4$ .  $\text{CH}_2$  is its empirical formula.

## Did you know?

The Galaxy Opal is the largest polished opal in the world. Opals consist mainly of silicon dioxide.



**C**

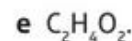
All substances have an **empirical formula**. This is the simplest whole number *ratio* of atoms or ions of each element in a substance.

Substances that are made of simple molecules also have a **molecular formula**. This represents the *actual* number of atoms of each element in one molecule.

Ethene has the molecular formula  $\text{C}_2\text{H}_4$  but the empirical formula  $\text{CH}_2$ . Sometimes the empirical formula is the same as the molecular formula. For example, the formula for a molecule of water is  $\text{H}_2\text{O}$  and this cannot be simplified any further.

The 'formula' of a compound is its molecular formula if it is a simple molecule, or the empirical formula if it has a giant lattice structure.

**1** Deduce the empirical formula of:



## Relative formula mass

The **relative formula mass** ( $M_r$ ), of a substance is the sum of the relative atomic masses ( $A_r$ ) of all the atoms or ions in its formula. Relative atomic masses are given in the periodic table.

## Worked examples W1

Calculate the  $M_r$  of carbon dioxide ( $\text{CO}_2$ ).

$$\begin{aligned} &= A_r(\text{C}) + (2 \times A_r(\text{O})) \\ &= 12 + (2 \times 16) \end{aligned}$$

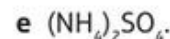
So,  $M_r$  of  $\text{CO}_2 = 44$

Calcium nitrate has a giant lattice structure. Its formula is  $\text{Ca}(\text{NO}_3)_2$  (for each calcium ion there are two nitrate ions). Calculate the  $M_r$  of calcium nitrate.

$$\begin{aligned} &= A_r(\text{Ca}) + 2(A_r(\text{N}) + (3 \times A_r(\text{O}))) \\ &= 40 + 2(14 + (3 \times 16)) \end{aligned}$$

So,  $M_r$  of  $\text{Ca}(\text{NO}_3)_2 = 164$

**2** Calculate the relative formula masses of:





## Finding an empirical formula

The empirical formula of a compound can be calculated from the masses of the elements used to make it. Table D shows how this is done if 10.0 g of calcium reacts with 17.8 g of chlorine.

Symbol for element	Ca	Cl
Mass (g)	10.0	17.8
Relative atomic mass, $A_r$	40	35.5
Divide the mass of each element by its relative atomic mass	$\frac{10.0}{40} = 0.25$	$\frac{17.8}{35.5} = 0.5$
Divide the answers by the smallest number to find the simplest ratio	$\frac{0.25}{0.25} = 1$	$\frac{0.5}{0.25} = 2$
Empirical formula	$\text{CaCl}_2$	

**D** To calculate an empirical formula, each element needs its own column of working.

A molecular formula is determined from its empirical formula and its  $M_r$ .

### Worked example W2

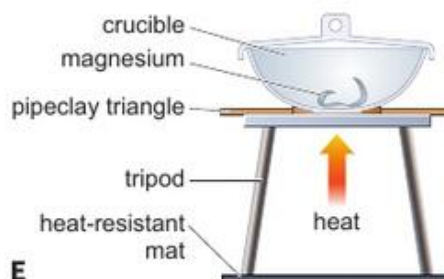
The empirical formula for glucose is  $\text{CH}_2\text{O}$  and its relative formula mass is 180. Determine the molecular formula for glucose.

- Find empirical formula mass  $A_r(\text{C}) + (2 \times A_r(\text{H})) + A_r(\text{O})$   
 $= 12 + (2 \times 1) + 16 = 30$
- Divide  $M_r$  by empirical formula mass  $\frac{180}{30} = 6$

The molecular formula is six times the empirical formula, so the molecular formula is  $\text{C}_6\text{H}_{12}\text{O}_6$ .

Magnesium oxide can be made by heating magnesium ribbon in a limited oxygen supply, using the apparatus in diagram E. If the reactant and product are 'weighed', the empirical formula for magnesium oxide can be calculated.

- 5** Magnesium ribbon with a mass of 0.576 g was heated in a crucible. It produced 0.960 g of magnesium oxide. Calculate the empirical formula of magnesium oxide.



- 3** Calculate the empirical formula of each of the following substances.



- a** lithium oxide  
(1.4 g Li, 1.6 g O)



- b** magnesium carbonate  
(1.2 g Mg, 0.6 g C, 2.4 g O)



- c** butane (1.44 g C, 0.3 g H)

- 4** Determine the molecular formula of each of the following substances.



- a** hydrogen peroxide, with empirical formula HO and  $M_r$  34



- b** hexene, with empirical formula  $\text{CH}_2$  and  $M_r$  84



- c** pentane, with empirical formula  $\text{C}_5\text{H}_{12}$  and  $M_r$  72

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** You are given some copper powder and a crucible. Describe an experiment to determine the empirical formula of copper oxide and explain how you would use the results to calculate the formula.

### Extend

- E1** A simple molecular compound contains 2.00 g of carbon, 0.33 g of hydrogen and 2.67 g of oxygen and has a relative formula mass of 60. Calculate the empirical and molecular formulae of this compound.

### Exam-style question

5.6 g of iron react with 24.0 g of bromine to form iron bromide. Calculate the empirical formula of the iron bromide. (Relative atomic masses are given in the periodic table at the back of the book.) (3 marks)

# SC9b Conservation of mass

Specification reference: C1.47; C1.48; C1.49

## Progression questions

- How do you calculate the concentration of a solution?
- How does the law of conservation of mass explain why magnesium increases in mass when it is burned?
- How do you calculate the masses of reactants and products in a reaction?

1 mg is 0.001 g and 1 l (1 litre) is the same volume as 1 dm<sup>3</sup>

Store in a cool dry place.  
Excerpt from the mineral water analysis by Laborunion Prof. Höll & Co. GmbH, Bad Elster dated April 01, 2004  
Composition of the characteristic ingredients:  
Typical values mg/litre:

Sodium	Na <sup>+</sup>	13.2
Calcium	Ca <sup>2+</sup>	29.1
Magnesium	Mg <sup>2+</sup>	3.0
Chloride	Cl <sup>-</sup>	31.1
Sulphate	SO <sub>4</sub> <sup>2-</sup>	42.7
Nitrate	NO <sub>3</sub> <sup>-</sup>	<0.5

The concentration of calcium ions is 29.1 mg/l or 0.0291 g dm<sup>-3</sup>.

**A** Mineral water contains dissolved ions.

## Concentrations of solutions

When a **solute** is dissolved in a **solvent** to make a **solution**, the mass of the solution is equal to the mass of the solvent *and* the mass of the solute. The overall mass of the substances does not change. This is the **law of conservation of mass**.

The amount of solute dissolved in a stated volume of a solution is its **concentration**. The units of concentration are usually 'grams per cubic decimetre', written as g dm<sup>-3</sup>. 1 dm<sup>3</sup> is the same volume as 1 litre or 1000 cm<sup>3</sup>.

You can calculate the concentration of a solution in g dm<sup>-3</sup> using this equation:

$$\text{concentration} = \frac{\text{mass of solute in g}}{\text{volume of solution in dm}^3}$$

If the volume of the solution is given in cm<sup>3</sup>, convert it to dm<sup>3</sup> by dividing by 1000.

**1** Calculate the concentration, in g dm<sup>-3</sup>, of the solute in these solutions.



**a** 20 g of sodium chloride in 2 dm<sup>3</sup> of solution



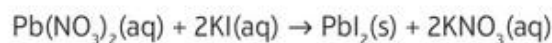
**b** 0.574 g of sodium hydroxide in 250 cm<sup>3</sup> of solution



**B** The total mass of the reactants always equals the total mass of products.

## Conservation of mass in reactions

Lead nitrate solution reacts with potassium iodide solution to form a yellow **precipitate** of lead iodide and a colourless solution of potassium nitrate. This is an example of a **closed system** as no new substances are added or removed.



The balanced equation shows that the number of atoms does not change, and so the mass cannot change. This is another example of the law of conservation of mass.

In the reaction shown in balanced equation C, the mass of the copper oxide left is less than the mass of copper carbonate at the start. Carbon dioxide gas escapes and so the mass appears to decrease. This is a **non-enclosed system** as the gas can escape.

## Did you know?

Lead iodide was used as a yellow pigment in some 19th-century paintings, but it is not very stable so some of these paintings are changing colour.



**C** balanced equation for heating copper carbonate in air



Some solids appear to gain in mass when they are heated in a non-enclosed system. In the reaction shown in balanced equation D, the apparent gain is due to copper reacting with oxygen from the air. However, in all systems, the mass of the products is always the same as the mass of the reactants.

## Calculating the masses of reactants or products

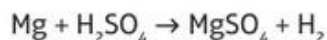
You can use relative masses and the balanced equation for a reaction to calculate the mass of a reactant or a product.

### Worked example

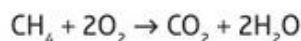
Calculate the mass of **chlorine** needed to make 53.4 g of **aluminium chloride**.

Write the balanced equation	$2\text{Al} + 3\text{Cl}_2 \rightarrow 2\text{AlCl}_3$
Calculate relative formula masses of the substances needed	$M_r \text{ Cl}_2 = 2 \times 35.5 = 71$ $M_r \text{ AlCl}_3 = 27 + (3 \times 35.5) = 133.5$
Calculate ratio of masses (multiply $M_r$ values by the balancing numbers shown in the equation).	
$3\text{Cl}_2$ makes $2\text{AlCl}_3$ so $3 \times 71 = 213 \text{ g Cl}_2$ makes $2 \times 133.5 = 267 \text{ g AlCl}_3$	
Work out the mass for 1 g of reactant or product. (Here we want 1 g of the product because that's the mass we know already.)	
$\div 267$	$\frac{213}{267} \text{ g Cl}_2$ makes $\frac{267}{267} \text{ g AlCl}_3$
	$0.798 \text{ g Cl}_2$ makes $1 \text{ g AlCl}_3$
$\times 53.4$	$42.6 \text{ g Cl}_2$ makes $53.4 \text{ g AlCl}_3$
Scale up or down (from 1 g to the mass you are given)	

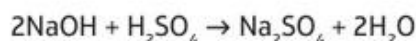
- 4** Calculate the mass of hydrogen produced when 72 g of magnesium reacts with sulfuric acid.



- 5** Calculate the mass of water produced when 500 g of methane burns.

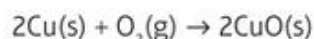
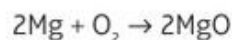


- 6** Calculate the mass of sodium hydroxide needed to produce 42.6 kg of sodium sulfate.



### Exam-style question

Calculate the maximum mass of magnesium oxide that could be formed by reacting 1.56 g of magnesium with excess oxygen. (3 marks)



**D** balanced equation for heating copper in air

- 2** 12.4 g of copper carbonate was heated and formed 8.0 g of copper oxide. Calculate the mass of carbon dioxide produced.
- 3** 1.27 g of copper was heated in air and formed 1.59 g of copper oxide. Calculate the mass of oxygen that reacted with the copper.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Make notes so that you can explain to a friend who missed the lesson:
- how to calculate the mass of sodium chloride formed when 5.3 g of sodium carbonate reacts with excess dilute hydrochloric acid  

$$\text{Na}_2\text{CO}_3 + 2\text{HCl} \rightarrow 2\text{NaCl} + \text{H}_2\text{O} + \text{CO}_2$$
  - why there is a loss in mass during the reaction.

### Extend

- E1** Calculate the loss in mass when 2.96 g of magnesium nitrate decomposes.
- $$2\text{Mg}(\text{NO}_3)_2(\text{s}) \rightarrow 2\text{MgO}(\text{s}) + 4\text{NO}_2(\text{g}) + \text{O}_2(\text{g})$$

# SC9c Moles

Specification reference: **H** C1.50; **H** C1.51; **H** C1.52; **H** C1.53

## Progression questions

- H** How do you calculate the number of moles and number of particles of a substance?
- H** What controls the mass of product formed in a reaction?
- H** How do you work out a balanced equation from the masses of reactants and/or products?

**H**



**A** Clockwise from top left: one mole (1 mol) each of iron(III) chloride, copper sulfate, potassium iodide, cobalt nitrate, potassium manganate(VII) and sodium chloride.

In everyday life we use special words to mean the amount of an item. A pair is two of that item and a dozen is 12 of them. In chemistry, a mole is 602 204 500 000 000 000 000 000 particles! This number is usually written in standard form as  $6.02 \times 10^{23}$  and is known as the **Avogadro constant**.

The SI unit symbol for 'mole' is **mol**. 1 mol of a substance contains the Avogadro constant number of particles. These can be atoms, molecules or ions and so you need to specify the type of particles.

The mass of one mole of a substance is the relative atomic mass ( $A_r$ ) or relative formula mass ( $M_r$ ) in grams. The relative atomic mass of magnesium is 24, so 1 mol of magnesium has a mass of 24 g and contains  $6.02 \times 10^{23}$  atoms.

You can calculate the number of moles of any substance using this equation:

$$\text{number of moles of substance} = \frac{\text{mass of substance (g)}}{A_r \text{ or } M_r}$$

**1** Give the mass of the Avogadro constant number of:



- a** carbon atoms      **b** sodium atoms.

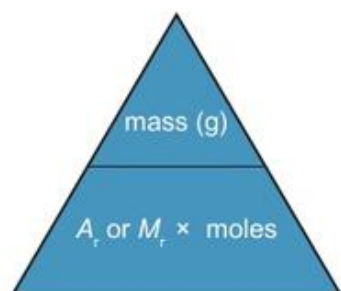
**2** Calculate the mass of:



- a** 2 mol of nitrogen molecules,  $N_2$



- b** 0.1 mol of sulfur dioxide molecules,  $SO_2$ .



**B** To rearrange the equation with this triangle, cover up the quantity you want to calculate and what you can see gives you the calculation to use.

**3** Calculate the number of moles of:



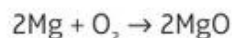
- a**  $CO_2$  molecules in 88 g of carbon dioxide



- b**  $CH_4$  molecules in 3.2 g of methane.

## Reactions

A balanced equation shows the ratios of the substances in moles.



This equation shows that 2 mol of magnesium reacts with 1 mol of oxygen to form 2 mol of magnesium oxide. In an experiment, we usually burn magnesium in excess oxygen to make sure that all the magnesium reacts.

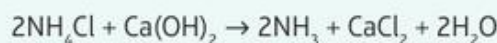
In a chemical reaction, one of the reactants is often added in excess and is not completely used up in the reaction. The amount of product formed is determined by the amount of reactant that is *not* in excess (and so is used up completely in the reaction). This is called the **limiting reactant**.



## H

## Worked example W1

1.50 g of ammonium chloride and 4.00 g of calcium hydroxide are heated together to form ammonia.



a Which is the limiting reactant?

b Calculate the mass of ammonia formed.

a The equation shows that 2 mol of  $\text{NH}_4\text{Cl}$  reacts with 1 mol of  $\text{Ca}(\text{OH})_2$   
 number of moles of  $\text{Ca}(\text{OH})_2 = 4.00 \text{ g} / (40 + 2(16 + 1)) = 0.0541 \text{ mol}$

We need:  $2 \times 0.0541 = 0.108 \text{ mol}$   $\text{NH}_4\text{Cl}$  to react with 0.0541 mol of  $\text{Ca}(\text{OH})_2$ .

We have:  $1.50 \text{ g} / (14 + (4 \times 1) + 35.5) = 0.0280 \text{ mol}$

We have less than the 0.0541 mol of  $\text{NH}_4\text{Cl}$  needed;  $\text{NH}_4\text{Cl}$  = limiting reactant.

b The equation shows that the number of moles of  $\text{NH}_3$  made equals the number of moles of  $\text{NH}_4\text{Cl}$  used.

So, 0.0280 mol of  $\text{NH}_4\text{Cl}$  forms 0.0280 mol of  $\text{NH}_3$

mass of  $\text{NH}_3$  formed = mol  $\times M_r = 0.0280 \times (14 + (3 \times 1)) = 0.476 \text{ g}$

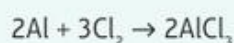
If you know the mass of each substance in a reaction, you can calculate the number of moles of each and deduce the balanced equation. The ratio of the moles of each substance is the **stoichiometry** of the reaction.

## Worked example W2

10.8 g of aluminium reacted with 42.6 g of chlorine,  $\text{Cl}_2$ , to produce aluminium chloride,  $\text{AlCl}_3$ . Deduce the balanced equation for the reaction.

	Al	$\text{Cl}_2$
Calculate the number of moles (= mass/ $A_r$ or $M_r$ )	$\frac{10.8}{27} = 0.4$	$\frac{42.6}{2 \times 35.5} = 0.6$
Divide by the smaller	$\frac{0.4}{0.4} = 1$	$\frac{0.6}{0.4} = 1.5$
Simplest whole number ratio	$1 \times 2 = 2$	$1.5 \times 2 = 3$

So 2 mol of Al react with 3 mol of  $\text{Cl}_2$ . The equation is completed by adding the formula of the product and balancing in the normal way.



## Exam-style question

Calculate the number of molecules in 90 g of water,  $\text{H}_2\text{O}$ .

(2 marks)

4 Calculate the number of molecules in:



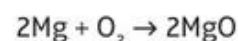
a 16 g of oxygen molecules,  $\text{O}_2$



b 34 g of ammonia molecules,  $\text{NH}_3$ .



5 Calculate the maximum mass of magnesium oxide that can be made from 2.4 g of magnesium and 2.4 g of oxygen.



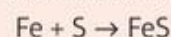
6 15 g of hydrogen gas reacts exactly with 70 g of nitrogen gas to produce ammonia,  $\text{NH}_3$ . Deduce the balanced equation for the reaction.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

S1 5.00 g of iron and 5.00 g of sulfur are heated together to form iron(II) sulfide.



a Calculate the number of moles of each reactant and state which one is the limiting reactant.

b Calculate the mass of iron(II) sulfide formed. Give your answer to 3 significant figures.

## Extend

E1 Explain which of these contains the greatest number of particles.

4 g of hydrogen molecules,  
 16 g of oxygen molecules, 18 g  
 of water molecules, 22 g of  
 carbon dioxide molecules.

## Empirical formula

A student carried out an experiment to determine the empirical formula of magnesium oxide using the apparatus shown.

Describe how you would use this apparatus and the results to show that the empirical formula of magnesium oxide is  $\text{MgO}$ .

(6 marks)



### Student answer

Weigh an empty crucible, then add a piece of magnesium ribbon and weigh it again. Set up the apparatus as shown. Heat the crucible with a strong flame and lift the lid occasionally [1]. Stop heating when all the magnesium has reacted. Let the crucible cool and reweigh it [2].

Find the mass of magnesium used and the mass of oxygen that combined with it [3]. Divide the masses of magnesium and oxygen by their relative atomic masses. Then, divide these answers by the smaller of the numbers to find the simplest ratio. If the simplest ratio is 1:1, the empirical formula is  $\text{MgO}$  [4].

[1] It would be a good idea to explain why this is necessary.

[2] Overall, this is a good account of the experiment and the measurements that are needed.

[3] The correct masses are mentioned, but not how these masses are actually calculated.

[4] The explanation of the calculation is sufficient and the relationship between the simplest ratio and the empirical formula is given.

### Verdict

This is a strong answer. The description of the experiment and the method of calculation are correct. The answer is written in a logical order with the description of the experiment first, and then a clear and well-ordered description of the calculations.

### Exam tip

The examiner is looking for a 'well-developed, sustained line of scientific reasoning which is clear and logically structured'. When describing a process, make sure that you write down the steps in a logical order – in this case, the order that the experiment and the calculations would be carried out.



# Paper 1

**SC10** Electrolytic

Processes /

**SC11** Obtaining and

Using Metals /

**SC12** Reversible

Reactions and

Equilibria /

**SC13** Transition Metals,  
Alloys and Corrosion

Many metals are used for buildings, including copper, aluminium and iron, together with alloys of those metals and others. In this unit you will learn more about metals.

## The learning journey

Previously you will have learnt at KS3:

- about oxidation and displacement reactions
- about the reactivity series.

You will also have learnt in *SC4 The Periodic Table*, *SC5 Ionic Bonding* and *SC8 Acids and Alkalis*:

- about anions and cations in ionic compounds
- to write balanced equations with state symbols
- how elements are arranged in the periodic table.

In this unit you will learn:

- more about reactivity, oxidation and reduction
- about how metals can be extracted
- about the advantages of recycling metals
- about the life cycle assessment of products
- to explain what happens during electrolysis and electroplating
- about equilibria in chemical reactions
- about the Haber process
- **H** how to write half equations
- about the properties of transition metals
- properties and uses of metals and their alloys.



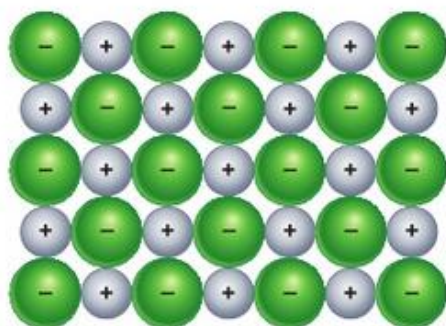


# SC10a Electrolysis

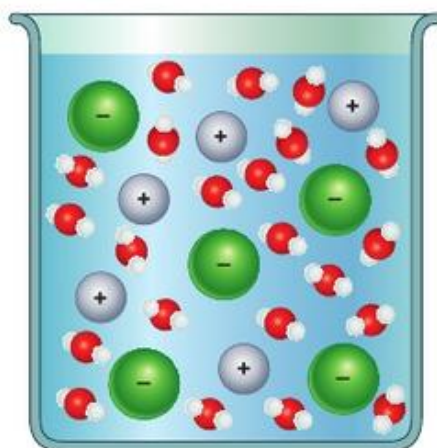
Specification reference: C3.22; C3.23; C3.24; **H** C3.27; **H** C3.28; **H** C3.29

## Progression questions

- What is an electrolyte?
- What happens to the ions during electrolysis?
- **H** How do you explain and represent the reactions taking place at the electrodes in electrolysis?



**A** (a) The ions cannot move in the lattice structure of solid sodium chloride.



(b) The ions can move when sodium chloride is dissolved in water.

In 1800, Alessandro Volta invented the electric battery. English scientists William Nicholson and Anthony Carlisle soon discovered that electricity broke down acidified water into hydrogen and oxygen. This process is called **electrolysis** and today we use it to break down many substances.

When an ionic solid is melted or dissolved in water, its ionic bonds break. This allows the ions to move. An ionic substance with freely moving ions is called an **electrolyte** and can conduct electricity.

- 1** Give the meaning of the following terms.



**a** electrolysis

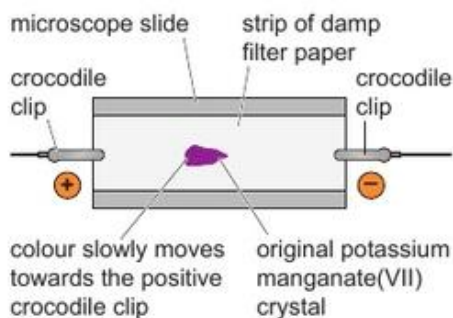


**b** electrode

Electrolysis uses energy transferred by electricity to decompose electrolytes. Two **electrodes** are connected to a direct current (d.c.) electricity supply and placed into the electrolyte. The two types of ions carry opposite charges and so migrate (move) towards the electrode with the opposite charge.

- **Cations** are positive ions and are attracted to the negative **cathode**.
- **Anions** are negative ions and are attracted to the positive **anode**.

This can be shown by placing a purple potassium manganate(VII) crystal on a piece of damp filter paper attached to a microscope slide and connected to a d.c. electricity supply. The purple colour spreads towards the anode.



**B** Manganate(VII) ions are purple. Potassium ions are colourless.



- 2** Give a reason why ionic solids cannot be electrolysed.



- 3** State the type of charge on a manganate(VII) ion, giving a reason for your answer.

- 4** Look at these the formulae:  $\text{Na}^+$ ,  $\text{Cl}^-$ ,  $\text{H}_2\text{O}$ ,  $\text{OH}^-$ ,  $\text{CO}_2$ ,  $\text{SO}_4^{2-}$ ,  $\text{Mg}^{2+}$



- a** Identify the anions.



- b** List the ions that will be attracted to the cathode during electrolysis.



## H Reactions at the electrodes

At the anode, negative ions lose electrons (**oxidation**).  
At the cathode, electrons are transferred from the electrode to the positive ions (**reduction**).

**Oxidation** Is the Loss of electrons.

**Reduction** Is the Gain of electrons.

You can remember these by: OIL RIG.

This transfer of electrons changes charged ions into atoms or molecules, resulting in chemical changes at the electrodes. For example, in the electrolysis of molten zinc chloride,  $\text{Zn}^{2+}$  ions are attracted to the cathode, where they gain electrons and become zinc atoms. At the same time  $\text{Cl}^-$  ions migrate to the anode, where they lose electrons and become chlorine molecules.

These changes are represented by **half equations**, which show the change at each electrode.

Cathode reaction:  $\text{Zn}^{2+} + 2\text{e}^- \rightarrow \text{Zn}$  reduction

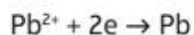
Redu**C**tion takes place at the **C**athode.

Anode reaction:  $2\text{Cl}^- \rightarrow \text{Cl}_2 + 2\text{e}^-$  oxidation

Oxid**A**tion takes place at the **A**node.

Note that two  $\text{Cl}^-$  ions are needed to form one chlorine molecule.

- 5 The half equation for the reaction at the cathode during the electrolysis of molten lead bromide is:



- a Explain whether this half equation shows oxidation or reduction.



- b Write the half equation for the reaction of bromide ions,  $\text{Br}^-$ , at the anode.



- 6 The following reaction takes place at an electrode:



Identify the electrode at which the reaction occurs, giving a reason for your answer.

## Did you know?

Metallic objects can be gold-plated using electrolysis. The anode is a strip of gold, the cathode is the object to be plated and the electrolyte contains gold ions.



C a gold-plated car

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1 Give the meaning of each of the terms: electrolyte, anode, cathode, anion, cation.  
S2 For each of the following ions, state which electrode it will move towards, during electrolysis:  $\text{Mg}^{2+}$ ,  $\text{OH}^-$ . Give a reason for your answers.

## Extend

- E1 **H** Molten potassium bromide is electrolysed to form potassium and bromine. Write the half equations for the reactions at the electrodes, classify them as oxidation or reduction and state at which electrode they occur.

## Exam-style question

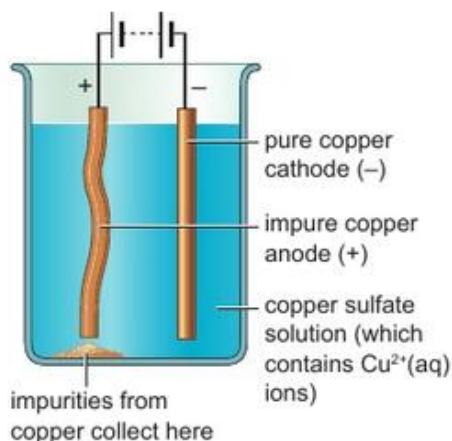
Solid lithium chloride cannot be electrolysed, but molten lithium chloride can be electrolysed. Explain these observations. (2 marks)

# SC10a Core practical – Electrolysis of copper sulfate solution

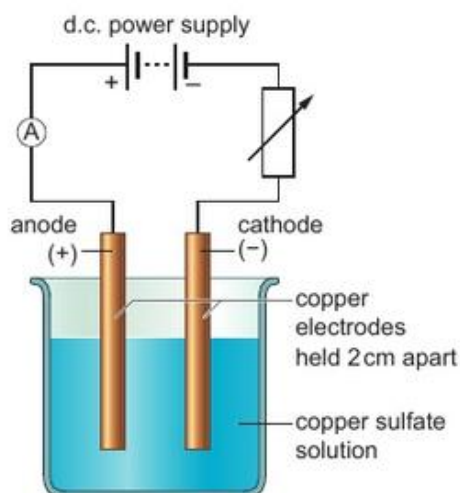
Specification reference: C3.31

## Aim

Investigate the electrolysis of copper sulfate solution with inert electrodes and copper electrodes.



**A** In the purification of copper, the copper to be purified is used as the anode and some very pure copper is used as the cathode.



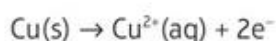
**B** Purifying copper by electrolysis. The copper to be purified is used as the anode and some very pure copper is used as the cathode.

The copper needed for electrical wires must be very pure and this is achieved using the electrolysis of copper sulfate solution, as shown in figure A.

During electrolysis, the copper atoms in the anode lose two electrons each to become copper ions. These ions dissolve in the solution and migrate to the cathode, where they are deposited as pure copper. So, the impure copper anode loses mass and the pure copper cathode gains mass. Impurities from the anode do not form ions and collect below the anode as a 'sludge'. The anode sludge is collected because it may contain valuable metallic elements.

## H

The half equation for the anode reaction is:



The half equation for the cathode reaction is:



## Your task

You will set up an electrolysis cell to investigate the effect of changing the current on the mass of the copper electrodes used in the electrolysis of copper sulfate solution. You will also investigate the products formed during the electrolysis of copper sulfate solution using inert (graphite) electrodes.

## Method

### Using copper electrodes

Wear eye protection.

- Select two clean pieces of copper foil. Label one 'anode' and the other 'cathode'. Measure and record the masses of the two electrodes.
- Set up an electrolysis circuit as shown in diagram B.
- Turn on the power and adjust the variable resistor to give a current of about 0.2 A. Record the current and adjust the variable resistor to keep it constant. Leave the power on for 20 minutes.
- Turn off the power and remove the electrodes from the beaker. Gently wash the electrodes with distilled water then dip them into propanone. Lift the electrodes out and gently shake off the propanone. Allow the remainder of the propanone to evaporate.
- Measure and record the masses of the dry electrodes.
- Repeat the experiment using currents of 0.3 A, 0.4 A and 0.5 A.

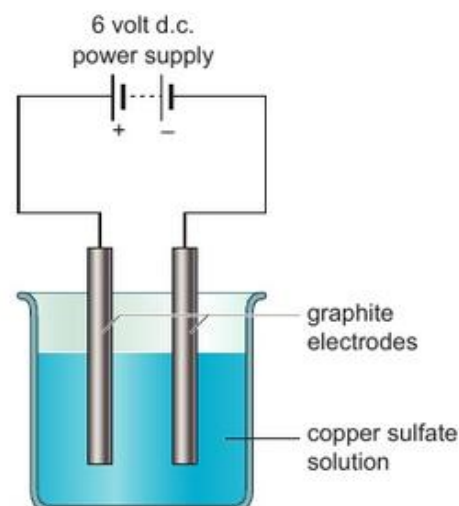


## Using graphite electrodes

- G** Set up the circuit as shown in diagram C.  
**H** Turn on the power and observe what happens at each electrode.

## Exam-style questions

- 1** Explain why a different product is formed at the anode when copper sulfate solution is electrolysed using graphite electrodes rather than copper electrodes. (4 marks)
- 2** Look at the method for electrolysis using copper electrodes.
  - a** State and explain *one* safety precaution (1 mark)
  - b** State why it is important to use clean copper electrodes. (1 mark)
  - c** Give a reason why a variable resistor is used in the electrolysis circuit. (1 mark)
  - d** Suggest a reason why the electrodes are washed at the end of the electrolysis. (1 mark)
  - e** Suggest a reason why propanone is used after washing the electrodes with distilled water. (1 mark)
- 3** The results of an investigation of the electrolysis of copper sulfate solution using copper electrodes are given in table D.
  - a** Calculate the changes in mass of the electrodes. (2 marks)
  - b** Plot a suitable graph to look for a correlation between the change in mass of each electrode and the current. (4 marks)
  - c** Describe the pattern in the change in mass at each electrode. (2 marks)
  - d** Explain the changes in mass of each electrode. (4 marks)
  - e** Explain the effect of increasing the current on these changes in mass. (2 marks)
  - f** Predict the change in mass at the anode when the current is 0.35 A. (1 mark)
  - g** Suggest a reason why the change in mass at the cathode is not the same as the change in mass at the anode when the same current is used. (1 mark)
  - h** Describe how you could improve the experiment to obtain more accurate results at the cathode. (1 mark)



**C** electrolysis circuit for using graphite electrodes

Current (A)	0.2	0.3	0.4	0.5
Mass of anode at start (g)	2.77	2.68	2.53	2.36
Mass of anode at end (g)	2.69	2.55	2.36	2.15
Mass of cathode at start (g)	2.51	2.55	2.62	2.70
Mass of cathode at end (g)	2.58	2.66	2.76	2.87

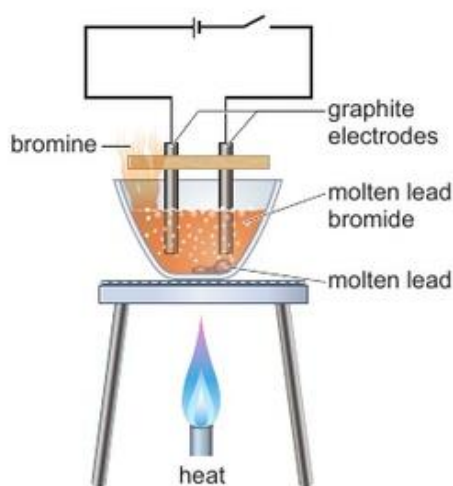
**D** results of an electrolysis investigation

# SC10b Products from electrolysis

Specification reference: C3.25; C3.26; C3.30

## Progression questions

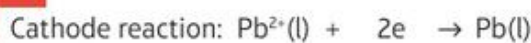
- How do you predict the products formed in the electrolysis of molten zinc chloride?
- How do you explain the products formed in the electrolysis of sodium chloride solution?
- How is copper purified using electrolysis?



**A** Orange bromine gas is seen around the anode and a pool of grey liquid lead is seen under the cathode. This experiment must be carried out in a fume cupboard as bromine gas is toxic.

The electrolysis of molten or dissolved ionic salts is carried out using **inert** (unreactive) electrodes (usually graphite or platinum). When a molten salt is electrolysed, ions are **discharged** as atoms or molecules at the electrodes. When molten lead bromide is electrolysed, bromine is produced at the anode and lead is produced at the cathode.

**H**



You can predict the electrolysis products of any molten salt. The salt will always decompose into its elements. The metal is produced at the cathode and the non-metal is produced at the anode.



- 1** Predict the products formed at the anode and cathode during the electrolysis of molten sodium chloride.



- 2** **H** Explain which of the half equations in the electrolysis of molten lead bromide shows oxidation and which shows reduction.

## Electrolysis of salt solutions

Water ionises to a very small extent, so in an aqueous solution of a salt there are some hydrogen ions ( $\text{H}^{+}$ ) and hydroxide ions ( $\text{OH}^{-}$ ), as well as the ions of the dissolved solid. Tables B and C show the electrolysis of two salt solutions, copper chloride solution and sodium chloride solution.

Ions	$\text{Cu}^{2+}(\text{aq})$ and $\text{Cl}^{-}(\text{aq})$ (from salt), $\text{H}^{+}(\text{aq})$ and $\text{OH}^{-}(\text{aq})$ (from water)	
cathode	$\text{Cu}^{2+}(\text{aq})$ and $\text{H}^{+}(\text{aq})$ ions are attracted. Copper ions are discharged more readily than hydrogen ions, so copper is formed as a brown solid.	<b>H</b> $\text{Cu}^{2+}(\text{aq}) + 2\text{e}^{-} \rightarrow \text{Cu}(\text{s})$ (reduction)
anode	$\text{Cl}^{-}(\text{aq})$ and $\text{OH}^{-}(\text{aq})$ ions are attracted. Chloride ions are discharged more readily than hydroxide ions, so chlorine is formed as a pale green gas.	<b>H</b> $2\text{Cl}^{-}(\text{aq}) \rightarrow \text{Cl}_2(\text{g}) + 2\text{e}^{-}$ (oxidation)
Overall	The copper chloride decomposes but the water does not change. $\text{CuCl}_2(\text{aq}) \rightarrow \text{Cu}(\text{s}) + \text{Cl}_2(\text{g})$	



- 3** Describe what happens to the colour of the solution when copper chloride is electrolysed.

**B** copper chloride solution electrolysis



Ions	Na <sup>+</sup> (aq) and Cl <sup>-</sup> (aq) (from salt), H <sup>+</sup> (aq) and OH <sup>-</sup> (aq) (from water)	
<b>cathode</b>	Na <sup>+</sup> (aq) and H <sup>+</sup> (aq) ions are attracted. Hydrogen ions are discharged more readily than sodium ions, so hydrogen gas is formed.	$\text{H} \quad 2\text{H}^+(\text{aq}) + 2\text{e}^- \rightarrow \text{H}_2(\text{g})$ (reduction)
<b>anode</b>	Cl <sup>-</sup> (aq) and OH <sup>-</sup> (aq) ions are attracted. Chloride ions are discharged more readily than hydroxide ions, so chlorine is formed as a pale green gas.	$\text{H} \quad 2\text{Cl}^-(\text{aq}) \rightarrow \text{Cl}_2(\text{g}) + 2\text{e}^-$ (oxidation)
<b>Overall</b>	The sodium chloride decomposes to form hydrogen and chlorine. The sodium and hydroxide ions remain in the solution. $2\text{NaCl}(\text{aq}) + 2\text{H}_2\text{O}(\text{l}) \rightarrow \text{H}_2(\text{g}) + \text{Cl}_2(\text{g}) + 2\text{NaOH}(\text{aq})$	

**C** sodium chloride solution electrolysis

In the electrolysis of sodium sulfate solution, sodium ions (Na<sup>+</sup>) and hydrogen ions (H<sup>+</sup>) collect at the cathode, where hydrogen gas forms. At the anode, hydroxide ions (OH<sup>-</sup>) are discharged more readily than sulfate ions (SO<sub>4</sub><sup>2-</sup>), so oxygen gas is formed (along with water). Overall, water decomposes to form hydrogen and oxygen. The Na<sup>+</sup> and SO<sub>4</sub><sup>2-</sup> ions stay in solution.

Water decomposes to form hydrogen and oxygen during the electrolysis of water acidified with dilute sulfuric acid (which contains H<sup>+</sup>(aq) and SO<sub>4</sub><sup>2-</sup>(aq)).

- 5** Use table C as a template to show the details of what happens in the electrolysis of:



**a** sodium sulfate

**b** acidified water.



- 6** Explain why the electrolysis of acidified water produces twice as much hydrogen as oxygen.

Copper can be purified by the electrolysis of copper sulfate solution using copper electrodes. The copper atoms in the anode lose electrons to become copper ions. These dissolve in the solution and migrate to the cathode, where they are deposited as pure copper. Impurities from the anode do not form ions and collect below the anode as a 'sludge'.



- 7**  $\text{H}$  Write the half equations for the reactions at the electrodes when copper sulfate is electrolysed using copper electrodes and classify them as oxidation or reduction.

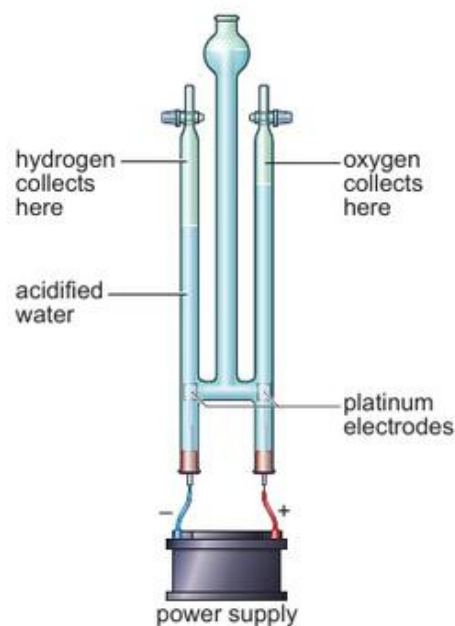
**Exam-style question**

The electrolysis of sodium chloride solution does not produce metallic sodium. State how you would change the electrolyte to obtain metallic sodium.

(1 mark)



- 4** When sodium chloride solution is electrolysed, state why hydrogen is produced instead of sodium.



**D** Water is electrolysed in a Hofmann voltameter to collect the gases produced.

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

- S1** Explain the formation of the products during the electrolysis of molten zinc chloride.
- S2** Predict the product formed at each electrode during the electrolysis of molten magnesium bromide. Justify your answers.

**Extend**

- E1** Explain why the electrolysis of sodium chloride solution produces hydrogen and chlorine at the electrodes.

# SC11a Reactivity

Specification reference: C4.1; **H** C4.2; C4.3

## Progression questions

- What are the similarities and differences in the way different metals react with water, acids and salt solutions?
- What happens to metal atoms when they react with water and acids?
- **H** How do you explain displacement reactions as redox reactions?




**A** Caesium is more reactive than potassium, and so is placed above potassium in the reactivity series.

### Did you know?

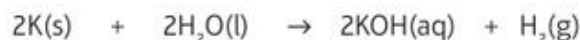
Caesium forms cations so easily that scientists think it could be used as a propellant in 'ion engines' for spacecraft.

The **reactivity series** is a list of metals in order of reactivity, with the most reactive at the top.

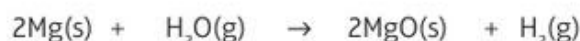
Metal	Reaction with water	Reaction with dilute acid	Tendency of metal atoms to form cations
potassium	react with cold water to form hydrogen and a metal hydroxide	react violently	
sodium			
calcium		react to form hydrogen and a salt solution	
magnesium	react very slowly, if at all, with cold water but react with steam to form hydrogen and a metal oxide		
aluminium			
zinc			
iron			
copper	do not react with cold water or steam	do not react	
silver			
gold			

**B** the reactivity series for some metals

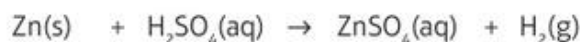
The metals that react with cold water form hydrogen and a metal hydroxide solution. For example:



The metals that react with steam form hydrogen and a solid metal oxide.



The metals that react with dilute acids form hydrogen and a salt solution. Bubbles of gas will be seen. The more bubbles formed, the more reactive the metal.



In all of these reactions, the metal atoms lose electrons to form positive ions (**cations**). The more easily a metal's atoms lose electrons, the higher the metal is in the reactivity series.

**6th** 1 Name a metal that does not react with cold water or dilute acid.

**7th** 2 Write the word equation for the reaction of calcium with water.

**9th** 3 Write the balanced equation for the reaction of magnesium with dilute hydrochloric acid. Include state symbols.



## Displacement reactions

We can use the reactivity series to predict whether reactions will take place. Each metal will react with compounds of the metals below it in the series.

When zinc is dipped into copper sulfate solution, a copper coating forms on the surface of the zinc. Some of the zinc takes the place of the copper and forms zinc sulfate solution.



This is a **displacement reaction**. The zinc has displaced the copper.

Displacement reactions only work one way. Copper cannot displace zinc from its compounds because copper is less reactive.



- 4 Use the reactivity series to predict whether a displacement reaction will take place in the reaction below. Either complete the equation or write 'no reaction'.



### H

Displacement reactions are also redox reactions. The reaction between zinc and copper sulfate can be written as an ionic equation:



The sulfate ions have been left out as they do not change and are called **spectator ions**. The zinc atoms lose electrons to form zinc ions. This can be shown in a **half equation**. 2e represents the two electrons.



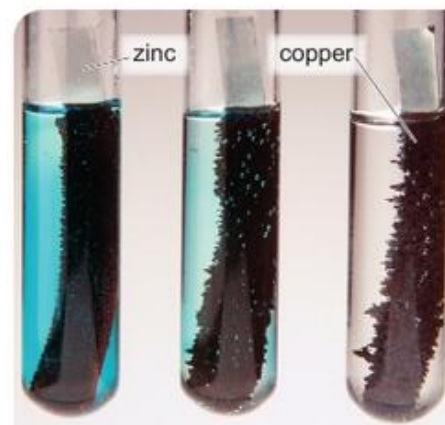
The copper ions gain electrons to form copper atoms:



A reaction in which one substance is oxidised and another is reduced is called a **redox reaction**.



- 6 Explain which substances are oxidised and reduced when:



C Blue copper sulfate solution changes to colourless zinc sulfate solution as zinc displaces copper.



- 5 Write the balanced equation for the reaction between magnesium and zinc chloride.

## Checkpoint

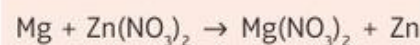
How confidently can you answer the Progression questions?

## Strengthen

- S1 Describe the reactions you could carry out to find the order of reactivity for magnesium, tin, chromium and copper. You may use the metals and the metal nitrate solutions.

## Extend

- E1 H Magnesium reacts with dilute sulfuric acid and zinc nitrate solution, as shown:



- Write ionic equations for these reactions.
- Explain, in terms of electrons, why these are redox reactions.

## Exam-style question

Three metals are labelled **X**, **Y** and **Z**. Metal **X** displaces **Z** from **Z** sulfate solution. Metal **X** does not displace **Y** from **Y** sulfate solution.

State the order of reactivity of the three metals, giving a reason for your answer. (3 marks)

# SC11b Ores

Specification reference: C4.4; C4.7; **H** C4.8

## Progression questions

- Which metals are found uncombined in the Earth's crust?
- How is the method of extraction of a metal related to its position in the reactivity series?
- **H** How are biological methods used to extract some metals?



**A** Gold occurs uncombined in the Earth's crust.

Very unreactive metals, such as gold and platinum, are found naturally in their **native state** (as uncombined elements). More reactive metals have reacted with other elements to form compounds in rocks. The process of obtaining a metal from these compounds is **extraction**.

An **ore** is a rock that contains enough of a compound to extract a metal for profit. Haematite is an ore containing iron oxide. Iron is extracted by heating the iron oxide with carbon. Carbon is more reactive than iron so it displaces it.



This method is used for compounds of metals below carbon in the reactivity series shown in table B.

- 7<sup>th</sup>** 1 Write word equations for the two reactions needed to produce copper.

- 7<sup>th</sup>** 2 State why copper can be produced by heating copper oxide with carbon.

Malachite is an ore containing copper carbonate. Malachite is heated to convert it to copper oxide, which is then heated with carbon to produce copper.

Metals higher than carbon must be extracted using a more powerful method called **electrolysis**. This involves passing electricity through a molten ionic compound to decompose it into its elements. Aluminium is produced by electrolysis of aluminium oxide, found in an ore called bauxite:



increasing reactivity	Metal	Method of extraction
	potassium	electrolysis of a molten compound
	sodium	
	calcium	
	magnesium	
	aluminium	
	(carbon)	
	zinc	heat an ore with carbon
	iron	
	copper	
	silver	found as the uncombined element
	gold	

A lot of energy is needed to keep metal oxides molten for electrolysis, making it extremely expensive. Electrolysis is only used to extract very reactive metals that cannot be obtained by heating their oxides with carbon.

- 5<sup>th</sup>** 3 Name two metals that can only be extracted from their ores by electrolysis.
- 4 Zinc can be extracted from zinc oxide, ZnO, by heating with carbon.
- 9<sup>th</sup>** a Write the balanced equation for this reaction.
- 9<sup>th</sup>** b Explain why electrolysis is not used in the large-scale extraction of zinc.

**B** reactivity of some metals



## H Biological methods of metal extraction

Copper is traditionally extracted by heating copper sulfide (producing copper and sulfur dioxide). However, copper ores are running out and so we need to extract copper from ores containing much smaller amounts of copper compounds.

**Bioleaching** uses bacteria grown on a low grade ore. The bacteria produce a solution containing copper ions, called a **leachate**. Copper is extracted from the leachate by displacement using scrap iron, then purified by electrolysis. This method can also be used for metals such as nickel, cobalt and zinc.

**Phytoextraction** involves growing plants that absorb metal compounds. The plants are burnt to form ash, from which the metal is extracted.

Process	Advantages	Disadvantages
both bioleaching and phytoextraction	no harmful gases (e.g. sulfur dioxide) are produced causes less damage to the landscape than mining conserves supplies of higher grade ores	very slow
bioleaching	does not require high temperatures	toxic substances and sulfuric acid can be produced by the process, and damage the environment
phytoextraction	can extract metals from contaminated soils	more expensive than mining some ores growing plants is dependent on weather conditions

**D** some advantages and disadvantages of bioleaching and phytoextraction



**5** Suggest why bioleaching and phytoextraction are very slow.



**6** Copper ions,  $\text{Cu}^{2+}$ , are produced in solution during bioleaching. Write an ionic equation for the displacement reaction between these copper ions and iron. Include state symbols.



**C** The tree used for phytoextraction contains a nickel-rich sap, which makes it green.

### Did you know?

When aluminium was first extracted by electrolysis in the late 19th century it was more expensive than gold.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Explain why iron is extracted from iron oxide using a different method to the one used to extract aluminium from aluminium oxide.

### Extend

**E1 H** Compare and contrast biological and non-biological methods of metal extraction.

### Exam-style question

Sodium is more reactive than aluminium. Describe a method to extract sodium from sodium chloride. Give a reason for your answer. (3 marks)



# SC11c Oxidation and reduction

Specification reference: **H** C4.2; C4.5; C4.6; C4.9

## Progression questions

- How do you explain oxidation and reduction in terms of oxygen?
- What types of reaction happen to ores when metals are extracted?
- How is the position of a metal in the reactivity series related to its resistance to corrosion?



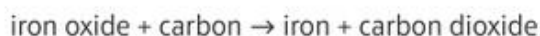
**A** Iron oxide is reduced to iron in a blast furnace.

Many metals are extracted from metal oxide ores. In order to obtain the metal from its oxide, the oxygen must be removed. When oxygen is removed from a compound, the compound is said to be 'reduced'.

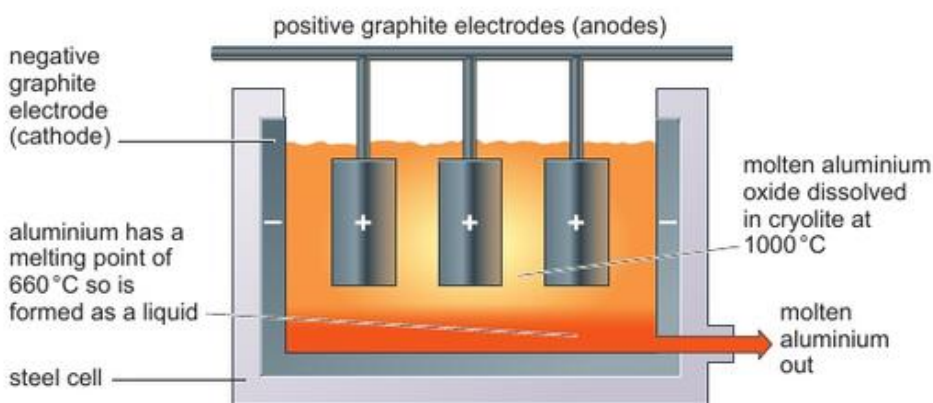
## Metal extraction is reduction

Oxidation is the gain of oxygen by a substance. Reduction is the loss of oxygen from a substance. Oxidation and reduction always occur together. If one substance is oxidised, another will be reduced. Reactions in which oxidation and reduction occur are called redox reactions.

Iron is obtained by removing the oxygen from iron oxide by heating with carbon. The iron oxide is reduced to iron. Carbon is oxidised to carbon dioxide.



Aluminium is obtained by removing oxygen from aluminium oxide by electrolysis.



**B** the electrolysis cell used to extract aluminium

During electrolysis, aluminium ions ( $\text{Al}^{3+}$ ) are attracted to the cathode, where they gain electrons to form aluminium. The oxide ions,  $\text{O}^{2-}$ , are attracted to the anodes, where they lose electrons to form oxygen. At the high temperature in the electrolysis cell, the oxygen reacts with the graphite (carbon) anodes to form carbon dioxide.

**H**

The half-equations for the reactions occurring at the electrodes are:



**1** Metals are obtained by reduction of their ores. State the meaning of 'reduction'.

**2** Zinc oxide reacts with carbon to form zinc and carbon dioxide in a redox reaction. Explain which substance has been oxidised and which has been reduced.

**3** **H** Molten sodium chloride contains  $\text{Na}^+$  and  $\text{Cl}^-$  ions. It is electrolysed to form sodium and chlorine.

**a** Write the half-equations for the reactions at the anode and the cathode.

**b** Explain whether the reaction at each electrode is oxidation or reduction.



## Corrosion

**Corrosion** happens when a metal reacts with oxygen, making the metal weaker over time. The metal gains oxygen so is oxidised. The corrosion of iron requires water as well as oxygen and is called **rusting**.

The more reactive a metal is, the more rapidly it corrodes. Gold does not corrode at all, which is one of the reasons it is used in jewellery.



**C** The Grouville Hoard was found in a field in Jersey in 2012. It contains about 70000 coins and pieces of gold jewellery, which are 2000 years old. The gold has not corroded.

Some metals, such as aluminium, are quite reactive and would be expected to corrode quickly. However, they do not corrode because their surfaces form a protective oxide layer (a **tarnish**), which prevents further reaction.



**D** The iron in the Angel of the North has reacted with oxygen and water.

### Exam-style question

Iron oxide is reduced by heating it with carbon. Explain what is oxidised in this process. (2 marks)

### Did you know?

The iron originally on Mars has rusted over billions of years, making the planet red. On the Moon, however, the dark areas on its surface are made of an iron-rich rock. The iron in these rocks has not rusted.



- 4** Look at the Did you know? box. Explain what the different colours in the iron-rich rocks on Mars and the Moon tell us about their atmospheres.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Copper oxide reacts with hydrogen:  
 $\text{CuO} + \text{H}_2 \rightarrow \text{Cu} + \text{H}_2\text{O}$   
 State, with a reason, which substance has been:  
**a** oxidised **b** reduced.
- S2** Three metals were left outside for 1 month. Metal A did not corrode at all; metal B corroded a lot; metal C corroded a little. Put the metals in order of increasing reactivity and explain your reasoning.

### Extend

- E1** **H** The half-equation for the reaction taking place at one of the electrodes during the electrolysis of molten magnesium chloride is:  
 $\text{Mg}^{2+} + 2\text{e}^- \rightarrow \text{Mg}$ . Explain: **a** at which electrode this reaction occurs **b** whether this is oxidation, reduction or neither.



# SC11d Life cycle assessment and recycling

Specification reference: C4.10; C4.11; C4.12

## Progression questions

- What are the advantages of recycling a metal?
- When might recycling a material not be worthwhile?
- What are the factors to consider in a life cycle assessment of a product?



**B** The WEEE man is made from over 3 tonnes of electrical and electronic equipment that the average person in the UK throws away in a lifetime.

- 1** Describe what is meant by 'recycling metals'.
- 2** Make a table to summarise the advantages and disadvantages of recycling metals compared to extracting more of the metals from ores.



**A** Artists have recycled the waste washed up on the coast of Kenya into sculptures.

Every UK household is estimated to produce 1 tonne of waste each year. About 45% of this waste is recycled but there is an EU target for **recycling** to reach 70% by 2030.

An estimated 2 million tonnes of waste electrical and electronic equipment (WEEE), such as phones, TVs and toasters, are thrown away in the UK each year. These items contain a lot of precious metals worth about £1 billion, including about £36 000 000 worth of aluminium.

## Recycling metals

Many metals can be recycled by melting them down and making them into something new. Some of the main advantages of recycling are as follows:

- Natural reserves of metal ores will last longer.
- The need to mine ores is reduced. Mining can damage the landscape as well as create noise and dust pollution.
- Less pollution may be produced. For example, sulfur dioxide is formed when some metals are extracted from metal sulfide ores.
- Many metals need less energy to recycle them than to extract new metal from the ore.
- Less waste metal ends up in landfill sites.

However, there are some disadvantages of recycling – including the costs and the energy used in collecting, transporting and sorting metals to be recycled. Sometimes it can be more expensive, and require more energy, to recycle than to extract new metal.



## Life cycle assessment

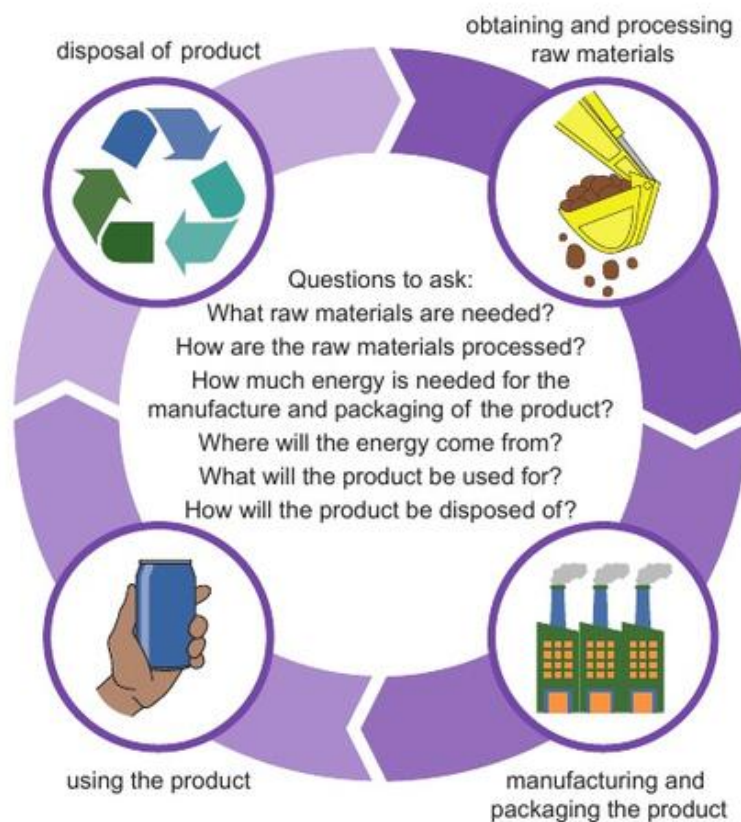
A life cycle assessment (LCA) can be carried out to work out the environmental impact of a product. An example is shown in diagram C.

The LCA also helps people to decide whether it is worthwhile to manufacture and recycle a product. LCAs can be used to compare the effect of using different materials for the same product, for example making a bottle from glass or plastic.

Some data that could be used as part of an LCA for the manufacture of a sample of 1 kg of aluminium is shown in the table below.

### Data on the manufacture of aluminium

- 5.5 kg of bauxite produces 1.9 kg of aluminium oxide, which produces 1.0 kg of aluminium during electrolysis
- 0.3 kg of carbon is burnt from the anodes
- 285 000 kJ of energy is needed to produce 1 kg of aluminium from bauxite
- 14 000 kJ of energy is needed to produce 1 kg of aluminium from recycled aluminium



C stages in an LCA

### Did you know?

The Colossus of Rhodes, one of the Seven Wonders of the Ancient World, was completed in 280 BCE using bronze from abandoned weapons. It was destroyed by an earthquake in 226 BCE. About 800 years later, the metal was recycled again.



D the Colossus of Rhodes



- 3 State the four stages that are considered in a life cycle assessment of a product.



- 4 Give two reasons why a life cycle assessment should be carried out before deciding whether to make a new product.

### Exam-style question

Describe two advantages of recycling copper, rather than obtaining it from its ores. (2 marks)

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1 a Outline how a broken aluminium saucepan is recycled to make an aluminium drinks can. b State the advantages and disadvantages of recycling aluminium.

### Extend

- E1 Discuss the life cycle assessment of aluminium, using the data shown in the data table and the extraction of aluminium in the previous topic. Include a description of other factors that need to be considered as part of the LCA.

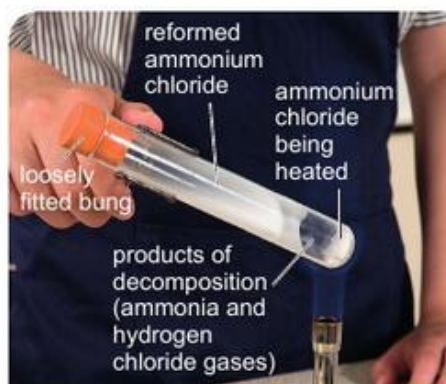


# SC12a Dynamic equilibrium

Specification reference: C4.13; C4.14; C4.15; C4.16; **H** C4.17

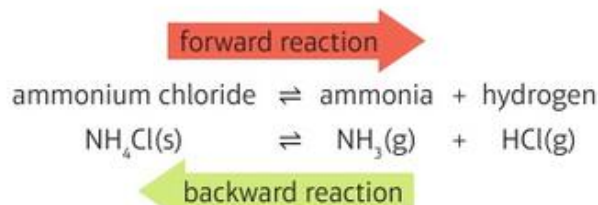
## Progression questions

- What is meant by dynamic equilibrium?
- How is ammonia manufactured?
- **H** How do changes in temperature, pressure and concentration affect the equilibrium position?



**A** Heating ammonium chloride is a reversible reaction.

In some chemical reactions the products react to reform the reactants. These are **reversible reactions**. In the equations for reversible reactions (such as the one shown in photo A), a double arrow ' $\rightleftharpoons$ ' is used to show that both forward and backward reactions occur at the same time.

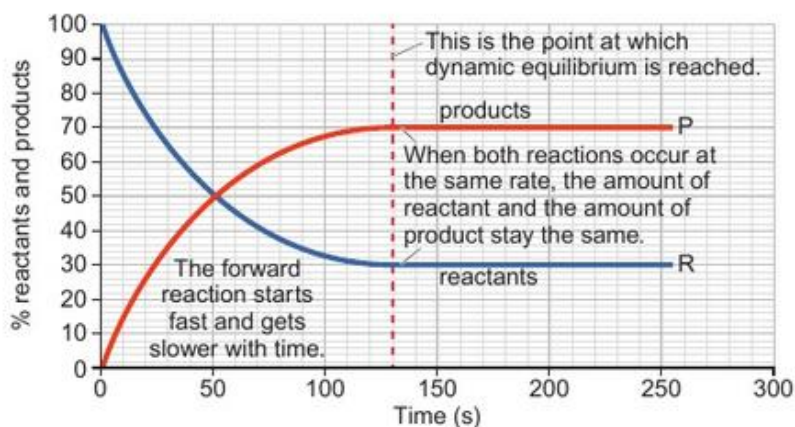


Graph B shows how the percentages of the reactants and products change during the reaction. At a certain point, the forward and backward reactions are still occurring but the percentages of the reactants and products are no longer changing. This is called a **dynamic equilibrium** because the reactions are still occurring (dynamic) but the substances remain in balance (equilibrium).

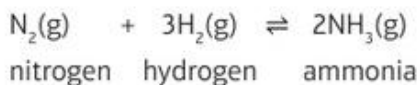
Although all reactions are reversible, dynamic equilibrium only occurs in **closed systems**, where there is no loss of reactants or products. In an **open system**, gases could escape and so equilibrium would not be achieved.

- 2** When calcium carbonate is heated in an open test tube it decomposes to form calcium oxide and carbon dioxide.
  - a** Write a word equation to represent this change as a reversible reaction.
  - b** Explain why equilibrium will not be achieved in this case.

- 7<sup>th</sup>** **1** What is happening when a reaction reaches 'dynamic equilibrium'?



**B** In a reversible reaction, the backward reaction gets faster with time, and the forward reaction gets slower with time. When they are occurring at the same rate, dynamic equilibrium has been reached.



The manufacture of ammonia by the Haber process involves a reversible reaction between nitrogen (from the air) and hydrogen (from natural gas) that can reach a dynamic equilibrium. The 'equilibrium position' (the percentages of the products and reactants at equilibrium) is changed by the reaction conditions. In the Haber process, and all similar industrial reactions, the reaction conditions are chosen to favour the forward reaction and make a large amount of product as cheaply as possible. In the Haber process, these conditions are a temperature of 450°C, a pressure of 200 atmospheres and the use of an iron catalyst.

- 8<sup>th</sup>** **3** What happens to the amount of ammonia after equilibrium is reached in the Haber process?



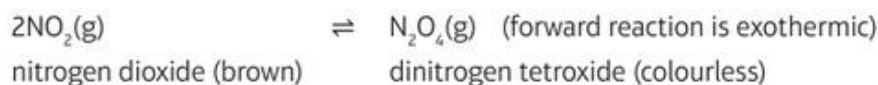
**H**

The equilibrium position can be altered by changes in temperature, pressure and concentration. In general, the equilibrium position shifts to reduce the effects of any changes to the system, as described in table C.

Change by ...	Equilibrium position shifts ...
increasing temperature	in the <b>endothermic</b> direction (transferring energy from the surroundings, cooling them down)
decreasing temperature	in the <b>exothermic</b> direction (transferring energy to the surroundings, heating them up)
increasing gas pressure	in the direction that forms fewer gas molecules (as this reduces pressure)
decreasing gas pressure	in the direction that forms more gas molecules (as this increases pressure)
increasing a concentration	in the direction that uses up the substance that has been added
decreasing a concentration	in the direction that forms more of the substance that has been removed

**C**

Photo D shows how the equilibrium position of a mixture of nitrogen dioxide and dinitrogen tetroxide depends on temperature.



**D** As the temperature is decreased the equilibrium shifts in the exothermic direction.

- 4** Explain why the equilibrium position in industrial reactions is often controlled.
- 5** Describe the difference between exothermic and endothermic reactions.
- 6** The formation of methanol from carbon monoxide and hydrogen is a reversible reaction that is exothermic in the forward direction.  

$$\text{CO}(\text{g}) + 2\text{H}_2(\text{g}) \rightleftharpoons \text{CH}_3\text{OH}(\text{g})$$

Explain the effect on the position of equilibrium of increasing:

- a** temperature
- b** pressure
- c** carbon monoxide concentration.

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

- S1** Use equations to explain the formation of ammonia in a reversible reaction that reaches a dynamic equilibrium.

**Extend**

- E1** Explain how a dynamic equilibrium is reached in the formation of ammonia.
- E2** **H** The formation of ammonia is exothermic. Describe three ways to increase the amount of ammonia at equilibrium.

**Exam-style question**

- a** Describe what happens in a reversible reaction. (1 mark)
- b** **H** Look at the equilibrium between  $\text{NO}_2$  and  $\text{N}_2\text{O}_4$  in diagram D. Explain why the colour changes when the pressure is increased. (3 marks)

# SC13a Transition metals

Specification reference: C5.1C

## Progression questions

- Where are the transition metals found in the periodic table?
- What are the typical properties of transition metals?
- What properties of iron make it a typical transition metal?



**A** Tungsten inert gas (TIG) welding joins pieces of steel together. The welding torch uses tungsten electrodes. The very high melting point of tungsten, 3422 °C, prevents these from melting.



- 1** Describe the position of the transition metals in the periodic table.



- 2** Name the property that allows steel car parts to be pressed into shape.



- 3** Bronze is a mixture of copper and tin. Suggest a reason that explains why some ancient Roman mirrors were made from polished bronze.

Tungsten and iron, like most other metals, are **transition metals**. These elements are placed in the central block of the periodic table, between groups 2 and 3. The transition metals include most of the metals used for construction, vehicles, electrical wiring, jewellery and other everyday uses.

1	2											3	4	5	6	7	0
			Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn						
								Rh	Pd	Ag							
					W			Ir	Pt	Au	Hg						

**B** the transition metals, with some common examples identified

## Physical properties

The transition metals have the typical properties of metals. In general, transition metals are:

- **malleable** (they can be hammered or rolled into shape without shattering)
- **ductile** (they can be stretched out to make thin wires)
- good conductors of electricity
- shiny when polished.

Iron is malleable and strong. It is used to make buildings, bridges, ships and cars. Copper is used for electrical wiring because it is ductile and a good conductor of electricity. Household mirrors consist of a sheet of glass coated with aluminium. Silver is even more shiny than aluminium, so it provides the reflective coating for the mirrors in infrared telescopes.

Compared to aluminium and the metals in groups 1 and 2, transition metals typically have:


- high melting points
- high densities.



Metal	Density (g/cm <sup>3</sup> )	Melting point (°C)
sodium	0.968	98
magnesium	1.74	650
aluminium	2.70	660
iron	7.87	1538
copper	8.96	1085
silver	10.5	962
tungsten	19.2	3422


**C** Sodium (group 1), magnesium (group 2) and aluminium (group 3) have lower densities and melting points than typical transition metals such as iron.

Remember that these are typical properties, and there are exceptions. For example, mercury is a transition metal with a high density, but it is liquid at room temperature.

-  **4** Explain why mercury is liquid, rather than solid or gas, at room temperature.


## Chemical properties

Aluminium, and the metals in groups 1 and 2, form white or colourless compounds. However, transition metal compounds are usually coloured. For example iron(III) oxide,  $\text{Fe}_2\text{O}_3$ , is red-brown. Tungsten oxide,  $\text{WO}_3$ , is yellow. Different transition metal hydroxides are different colours. This property is used to identify the transition metal ion present in a substance (see SC25b *Tests for positive ions*). Iron(II) hydroxide,  $\text{Fe}(\text{OH})_2$ , is pale green but iron(III) hydroxide,  $\text{Fe}(\text{OH})_3$ , is orange-brown.

-  **5** Rubies are red gemstones that are mostly aluminium oxide. Suggest a reason that explains why rubies are coloured.

Transition metals and their compounds typically show catalytic activity.

**Catalysts** increase the rate of a reaction without being changed either chemically or in mass at the end of the reaction. Iron is the catalyst used to manufacture ammonia by the Haber process. Iron(III) chloride,  $\text{FeCl}_3$ , is a catalyst used during the manufacture of poly(chloroethene) or PVC.

-  **6** Catalytic converters in vehicles convert harmful exhaust gases into less harmful ones. Suggest a reason that explains why these devices contain platinum, Pt, and rhodium, Rh.

## Exam-style question

Potassium, K, is a group 1 metal and titanium, Ti, is a transition metal. Predict *one* difference in their physical properties and *one* difference in their chemical properties. (2 marks)



**D** Mercury melts at  $-39^\circ\text{C}$  and boils at  $356^\circ\text{C}$ . A steel nail can float on it because of the high density of mercury,  $13.53\text{ g/cm}^3$ .



**E** Pure beryl is a colourless mineral. Iron impurities turn it into aquamarine, a blue gemstone.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** Why is iron considered to be a typical transition metal?

## Extend

**E1** Compare the typical properties of transition metals with those of other metals, such as sodium and aluminium.

# SC13b Corrosion

Specification reference: C5.2C; C5.3C

## Progression questions

- Why do metals corrode?
- How can the surface of iron be protected from rusting?
- How does sacrificial protection prevent rusting?

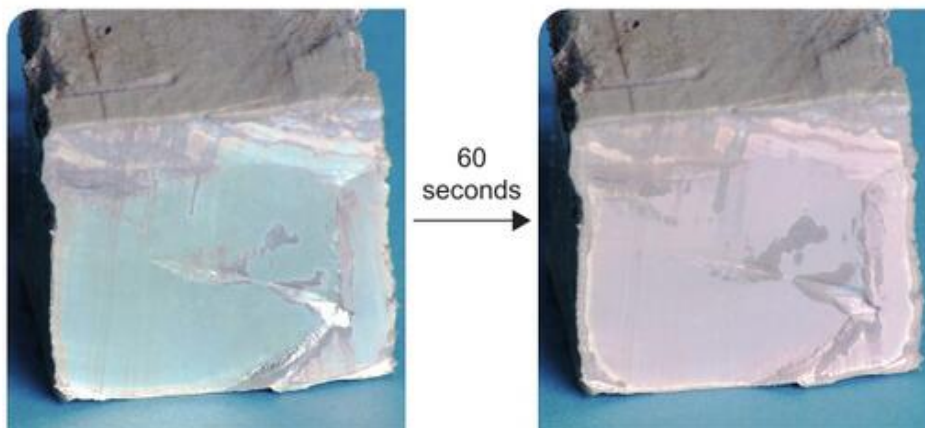
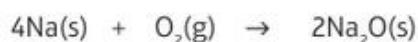


**A** The Statue of Liberty consists of 27 tonnes of copper sheeting over a steel frame. Its green colour is due to a natural layer of copper salts that forms when copper tarnishes.

- 6th** 1 Explain why magnesium ribbon appears dull grey, but becomes silvery when rubbed with sandpaper.
- 9th** 2 Write a balanced equation for the reaction of aluminium with oxygen, forming aluminium oxide,  $\text{Al}_2\text{O}_3$ .

When metals react with oxygen in the air, they **oxidise** to form metal oxides. Very reactive metals, such as sodium, oxidise rapidly when they are freshly cut:

sodium + oxygen  $\rightarrow$  sodium oxide



**B** White sodium oxide forms quickly on shiny, freshly cut sodium.

Less reactive metals, such as silver, oxidise slowly. Gold and other very unreactive metals may not react with oxygen at all.

Metals may form a thin layer of **tarnish** when they oxidise. This layer stops oxygen reaching the metal, preventing further oxidation. The green layer of 'verdigris' on the Statue of Liberty protects the copper underneath. In a similar way, aluminium is a reactive metal protected by a natural layer of its oxide. This layer is thin enough to see the aluminium underneath.

**H**

Metals oxidise when they lose electrons. This happens when metals react with oxygen. It may also happen when metals react with other substances found in air. For example, silver reacts with hydrogen sulfide to form a black layer of silver sulfide:

silver + hydrogen sulfide  $\rightarrow$  silver sulfide + hydrogen

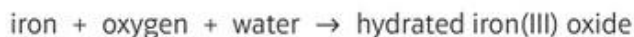


- 10th** 3 Write a half equation to model how silver oxidises to silver ions,  $\text{Ag}^+$ .



## Rusting

**Corrosion** happens when a metal continues to oxidise and so the metal becomes weaker over time. **Rusting** is the corrosion of iron or steel (which is mostly iron). Iron rusts when it reacts with oxygen and water:



Hydrated iron(III) oxide is the orange-brown substance seen on rusty objects. Rust flakes off rusty iron, which exposes more metal to air and water and so the iron continues to corrode and weaken.

**10** 4 Compare and contrast tarnishing with corrosion.

**9** 5 Explain why rusting is an oxidation reaction.

Rusting can be prevented if air is kept away. This can be achieved by storing the metal in an unreactive atmosphere of nitrogen or argon. Rusting can also be prevented if water is kept away, for example using a **desiccant** powder that absorbs water vapour. Other methods of rust prevention keep both air and water away, including painting, coating with plastic, oiling and greasing.

**6** 6 Explain why a bicycle chain is oiled rather than painted.

## Sacrificial protection

**Sacrificial protection** is a method of rust prevention that does not rely on keeping air or water away. Instead, a piece of magnesium or zinc is attached to the iron or steel object. Magnesium and zinc oxidise more easily than iron, so oxygen reacts with them rather than with the iron or steel object. This protection continues until the sacrificial metal corrodes away.

### H

The more reactive a metal is, the more easily it loses electrons. Zinc and magnesium lose electrons more easily than iron does, so they are more easily oxidised.



**D** blocks of sacrificial metal bolted onto a ship's hull

## Did you know?

The Iron Pillar of Delhi is a 6 tonne iron pillar, over 1600 years old and almost free of rust. It contains a relatively high amount of phosphorus. This forms a natural layer of iron hydrogen phosphate, which prevents air and water reaching the iron below.



**C** the Iron Pillar of Delhi

**8** 7 Explain why zinc blocks are attached to a ship's hull beneath its waterline.

## Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** How do different methods of rust prevention work?

### Extend

**E1** How could you investigate the effect of dissolved salt on the rate of rusting?

## Exam-style question

Evaluate the suitability of sodium for the sacrificial protection of an offshore oil rig.

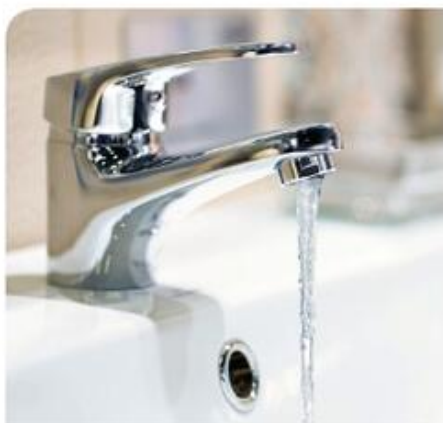
(3 marks)

# SC13c Electroplating

Specification reference: C5.4C

## Progression questions

- What is electroplating?
- Why are metals electroplated?
- How is electroplating done?



**A** Taps are plated with shiny chromium to improve their appearance and resistance to corrosion.



**B** High-end electrical connectors are plated with gold to improve their corrosion resistance.

## Did you know?

Replica ancient gold and silver coins are produced by electrotyping. The two sides of a real coin are pressed into lead to make a mould. The two moulds are electroplated with gold or silver. The electroplated metal halves are removed from the moulds, then filled and glued together to make the replica coin.

**Electroplating** coats the surface of one metal with a thin layer of another metal. Silver and gold are attractive transition metals but they are expensive. Silver or gold can be electroplated onto cheaper 'base metals' such as copper or nickel. This produces attractive jewellery that is cheaper than solid silver or gold.

Electroplating may also be used to improve a metal object's ability to resist corrosion. Chromium is a transition metal that resists corrosion. Objects such as vehicle parts and boat parts made from steel may be 'chrome plated' using electroplating. The thin layer of chromium stops air and water reaching the steel below, preventing the object rusting.



**1** Suggest two reasons that explain why a drum kit may be chrome plated.

Silver and copper are the best electrical conductors, but both tarnish in air. A layer of tarnish may form when cables and equipment are disconnected. Expensive hi-fi cables and electronic equipment may use gold-plated connectors. Gold is the third-best electrical conductor. It does not tarnish but it is much more expensive than silver or copper.

**2** Cables can be made from copper.



**a** Describe a disadvantage of using copper cables in electronic equipment.



**b** Give a reason that explains why gold-plated connectors are sometimes used in expensive hi-fi equipment.



**c** Describe a disadvantage of using gold-plated connectors in this type of equipment.

## Carrying out electroplating

To electroplate a metal object, you need:

- an **anode** (positively-charged electrode), made from the plating metal
- an **electrolyte**, which is a solution containing ions of the plating metal
- a **cathode** (negatively-charged electrode), which is the metal object itself.



**3** Describe three substances you need to electroplate a steel exhaust pipe with chromium.



Diagram C shows how you could electroplate a copper ring with silver. A direct current (d.c.) flows through the electrodes and the electrolyte. Silver ions in the electrolyte move to the negatively-charged copper ring. They gain electrons and are deposited as silver atoms. At the silver anode, silver atoms lose electrons to become silver ions, which go into the electrolyte. The longer the current flows, the thicker the silver layer on the ring becomes.

**H**

These half equations model the changes that happen when copper is electroplated with silver:

at the silver anode:  $\text{Ag(s)} \rightarrow \text{Ag}^+(\text{aq}) + \text{e}^-$

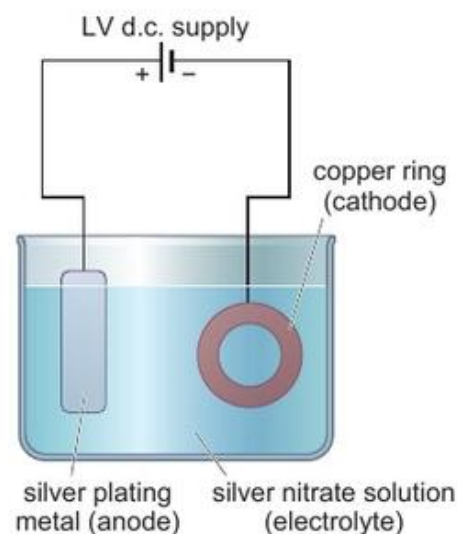
at the copper cathode:  $\text{Ag}^+(\text{aq}) + \text{e}^- \rightarrow \text{Ag(s)}$

8th

**4** Explain whether reduction or oxidation happens at the anode.

9th

**5** Metal bicycle parts may be electroplated with nickel to improve their appearance. Nickel chloride solution,  $\text{NiCl}_2(\text{aq})$ , or nickel sulfate solution,  $\text{NiSO}_4(\text{aq})$ , may be used as the electrolyte. Write half equations to model the changes that happen at each electrode.



**C** electroplating silver onto a copper ring

## Galvanising and tin plating

Iron and steel objects can be protected from rusting by coating them with zinc. This is called **galvanising**. The thin layer of zinc improves corrosion resistance by stopping the water reaching the iron or steel, and by acting as a sacrificial metal. This sacrificial protection can continue even if the zinc layer is damaged.

Galvanising can be carried out using electroplating, or by dipping the object in molten zinc.

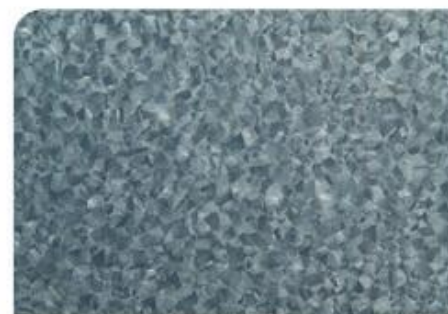
Food cans are made from steel. Their inner surface is protected from rusting by electroplating with tin. Tin does not react with air or water at room temperature. The tin layer stops air and water reaching the steel. However, if the tin layer is damaged, the steel can will rust faster. This is because iron is more reactive than tin, so it acts as a sacrificial metal to protect the tin.

8th

**6 a** Suggest reasons that explain why galvanising a steel farm gate is desirable.

7th

**b** Describe how you would use electroplating to galvanise the gate.



**D** the surface of galvanised steel

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Why are some metal objects electroplated with another metal?

### Extend

**E1** Evaluate the use of electroplating for jewellery and for metal bathroom fittings.

### Exam-style question

Suggest two reasons that explain why a gold medal may consist of silver electroplated with gold. (2 marks)

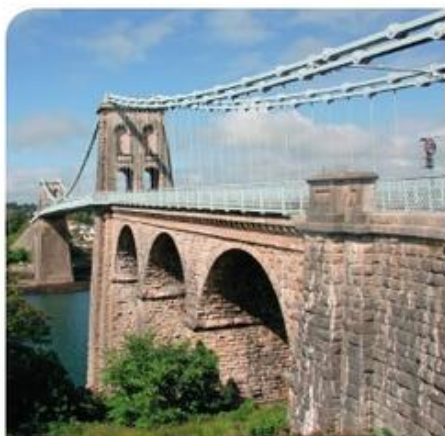


# SC13d Alloying




Specification reference: C5.5C; C5.6C

## Progression questions

- What is an alloy?
- Why is iron mixed with other metals to make alloy steels?
- Why are alloys often stronger than pure metals?



**A** The Menai Bridge was completed in 1826. Its original wrought iron suspension chains were replaced by steel chains in the 1930s to increase the bridge's weight limit.

-  **1** Explain why wrought iron is an alloy.
-  **2** Write a balanced equation for the reaction between chromium and oxygen.
-  **3** Suggest a reason that explains why stainless steel is used for surgical instruments.

## Did you know?

Alloys of iron, aluminium, carbon and manganese are lightweight but brittle. In 2015, Korean scientists discovered that adding some nickel produces an alloy as strong and lightweight as titanium, but costing ten times less.

An **alloy** is a mixture of a metal element with one or more other elements, usually metals. For example, iron straight from the blast furnace is not pure iron. It is an alloy of iron with about 4% carbon, and smaller proportions of phosphorus and other elements. Wrought iron is produced by lowering the carbon content to less than 0.08%.

## Alloy steels

**Alloy steels** are made by deliberately adding other elements to iron. It is possible to create a huge range of alloy steels with different properties by varying their composition.

**Stainless steels** resist rusting, unlike other alloy steels and iron. Stainless steels contain chromium, which reacts with oxygen in the air. A layer of chromium oxide,  $\text{Cr}_2\text{O}_3$ , forms. This is thick enough to stop air and water reaching the metal below, but thin enough to be transparent. If the metal is scratched, more chromium reacts to replace the layer.

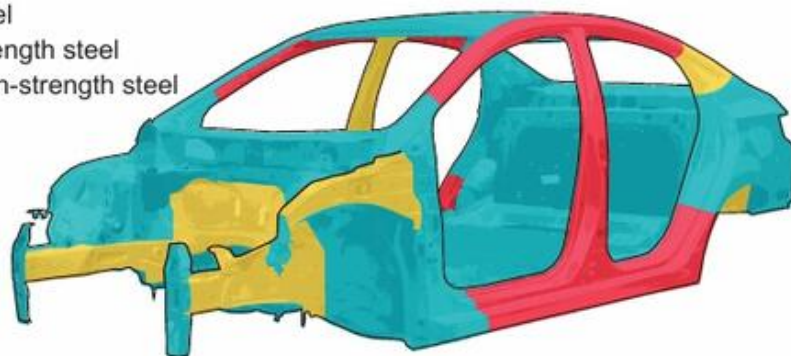


**B** Stainless steel is useful for domestic and professional kitchens.

Some alloy steels are very strong and tough. 'Tool steels' contain tungsten and molybdenum, and are used to make drill bits. In general, the greater the carbon content of steel, the stronger and harder it is. Mild steel has a low carbon content. However, it also contains elements such as manganese to increase its strength while maintaining its malleability. Mild steel is often used as a building material and for car body panels. Stronger steels can be more difficult to press into shape, but car manufacturers use them to produce strong but relatively lightweight car bodies.



- mild steel
- high-strength steel
- ultra high-strength steel

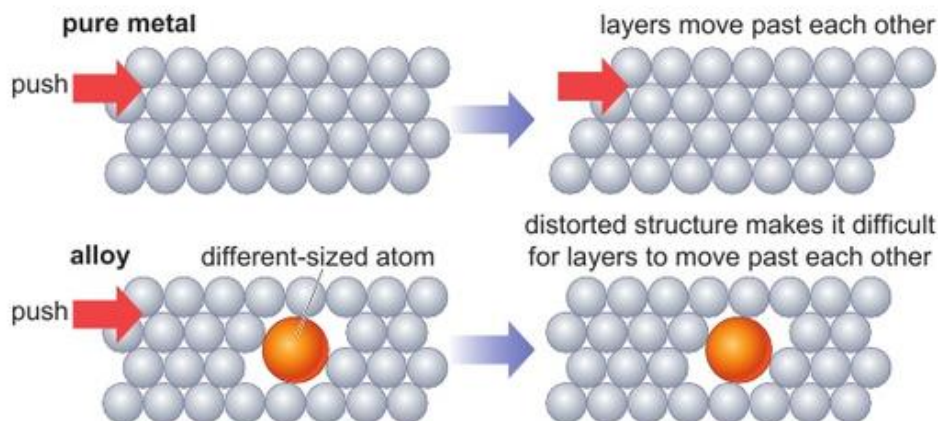


**C** Different alloy steels are used to make a car body.

## Alloys and strength

Alloys are often stronger than the pure metals they contain. For example, pure iron is too soft for everyday use, but wrought iron is stronger even though it contains only a small amount of other elements. Alloy steels are even stronger than wrought iron.

In a solid, pure metal the atoms are all the same size and are arranged regularly in layers. These layers move past each other if enough force is applied. This is why metals are malleable and ductile. In an alloy, the atoms of other elements present may be different sizes. They distort the regular structure, making it more difficult for the layers to slide past each other. This is why alloys are often stronger than pure metals, even though they are usually still malleable and ductile.



**D** explaining the difference in strength between pure metals and alloys

Diagram D shows a simple particle model to explain the difference in strength between a pure metal and an alloy. This is a two-dimensional model, but remember that metals have a three-dimensional structure.

- 6** Explain why car parts are made from alloy steels, rather than from pure iron or wrought iron.

- 4** A car body made from mild steel alone would be heavier than one that also contains high-strength steel and ultra high-strength steel (see diagram C). Give a reason that explains this.

- 5** Explain, in terms of their structure, why alloys are stronger in *all* directions than the pure metals they contain.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** Explain, with examples, why metals are often used as alloys rather than as pure metals.

## Extend

- E1** Evaluate the statement 'alloys are stronger than the individual metals they contain'.

## Exam-style question

Jewellery gold is often an alloy of gold and copper. Explain how the presence of copper atoms can produce an alloy stronger than pure gold. (3 marks)



# SC13e Uses of metals and their alloys

Specification reference: C5.7C

## Progression questions

- What are some common uses for aluminium, copper and gold?
- What are some common alloys containing aluminium or copper?
- Why are different metals or their alloys chosen for different uses?



**A** Magnalium is the name given to a range of alloys of magnesium and aluminium. When it has a high magnesium content, magnalium is brittle and flammable, making it suitable for use in signal flares and sparklers.

The uses of a metal or alloy depend upon its properties. These properties include:

- chemical properties, such as resistance to corrosion
- physical properties, such as density and ability to conduct electricity.

For example, gold and copper resist corrosion. They are also malleable, ductile and very good conductors of electricity. Both metals could be used for electrical wiring, but cost may also be an important factor. Gold is thousands of times more expensive than copper. So copper is chosen for most electrical wiring, while gold is used in tiny amounts to connect microprocessors and memory chips.



**B** This restorer is applying pieces of very thin gold, called gold leaf, to an antique ornament.

Aluminium resists corrosion, but it does not conduct electricity as well as copper. However, it is stronger, cheaper and less dense, so it is used for overhead electrical cables. The properties of a metal may also depend upon the intended use. For example, several wires can be wound around each other to make a cable. Such a cable is stronger than one made from a single wire of the same diameter.



**C** Aluminium is used in overhead electrical cables, such as these being fitted to the National Grid's new T-pylons.

-  **1** Suggest reasons that explain why copper is used to make plumbing parts such as water pipes.
-  **2** Suggest reasons that explain why very thin gold leaf is used to decorate the surfaces of statues and picture frames.





**D** Spacesuit visors are coated with a thin layer of gold. This reflects most of the Sun's harmful infrared radiation, but is thin enough to let visible light through.

Alloys may have more useful properties than the metals they contain. For example, brass is an alloy of copper and zinc. Copper and brass both resist corrosion. Copper is a better conductor of electricity than brass, but brass is stronger. This makes brass more suitable than copper for making electrical plug pins.

Magnalium containing 95% aluminium and 5% magnesium is an engineering alloy used for aircraft parts and scientific instruments. Table E shows some properties of the two pure metals and the alloy. Magnalium is less dense and almost four times stronger than aluminium alone. Although it is denser than magnesium, magnalium is twice as strong and has better resistance to corrosion. These properties allow the manufacture of strong but lightweight metal parts.

Metal	Density (g/cm <sup>3</sup> )	Relative strength
aluminium	2.70	1.0
magnesium	1.74	1.9
magnalium	2.50	3.8

**E** properties of aluminium, magnesium and 95% Al / 5% Mg magnalium

- 6** Explain, using information from table E, why engineering magnalium is more suitable for aircraft parts than aluminium or magnesium alone.

### Exam-style question

Jewellery gold is often an alloy of gold and copper. Other than increasing the strength of the jewellery, give two reasons why copper is chosen to mix with gold. (2 marks)

- 3** Suggest reasons that explain why some overhead electricity cables consist of steel wires with aluminium wires wound around them.
- 4** Evaluate the choice of metal, copper or aluminium, in household electricity cables and overhead electricity cables.

### Did you know?

Gold is incredibly malleable and ductile. Just 1 g of gold can be hammered into a 60 cm diameter circle only 0.18  $\mu\text{m}$  thick, or made into a wire nearly 3 km long.

- 5** Explain why brass is suitable for making coins and musical instruments.

### Checkpoint

How confidently can you answer the Progression questions?

#### Strengthen

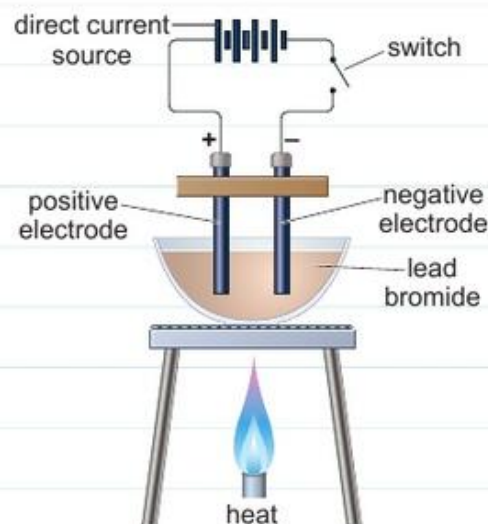
**S1** Why are different objects made from different metals?

#### Extend

**E1** Describe to what extent alloys are a better choice for a given purpose than a pure metal.

# Electrolysis

Lead bromide contains  $\text{Pb}^{2+}$  and  $\text{Br}^-$  ions and has the formula  $\text{PbBr}_2$ . The diagram shows the electrolysis of lead bromide. Explain what is happening at each electrode at the start of the experiment and as the electrolyte melts. (6 marks)



## Student answer

Lead bromide contains  $\text{Pb}^{2+}$  ions and  $\text{Br}^-$  ions and has the formula  $\text{PbBr}_2$  [1]. It is a compound and has ions, so it can be used for electrolysis [2]. During electrolysis, the ions move towards the oppositely-charged electrodes [3]. The products at each electrode are lead and bromine [4]. Heat is needed to melt the lead bromide so that the ions can move and conduct electricity. When the lead bromide is a solid, the ions can't move, so there is no flow of electricity.

[1] There are no marks for simply repeating information from the question.

[2] This is correct, but it is not relevant to the question which asks students to explain what is happening at the electrodes.

[3] The answer would be better if it explained what was happening at each electrode.

[4] These are the correct products, but the answer should state which product is formed at each electrode.

## Verdict

This is an acceptable answer. It contains some correct chemistry, but is missing some information. There is a good explanation of why heat is needed for electrolysis, and the products of the electrolysis are given.

The answer could be improved by explaining which ions move towards which electrode and what happens at those electrodes. For example, it could explain that the positively-charged cations ( $\text{Pb}^{2+}$ ) move towards the negatively-charged electrode (cathode), and once there, they gain two electrons to become lead. It would also have been more logical to have the last two sentences earlier in the explanation, starting with what happens in the circuit at the start of the experiment (when the lead bromide is still solid). There is also information that is copied from the question and some parts of the answer are not relevant.

## Exam tip

Do not just repeat information from the question, and make sure that anything you write is relevant to the question that has been asked.



# Paper 1

## SC14 Quantitative Analysis / SC15 Dynamic Equilibria, Calculations Involving Volumes of Gases / SC16 Chemical Cells and Fuel Cells

The photo shows an ethanol production facility, where sugars are fermented into ethanol in fermentation tanks. The ethanol is then purified for use as transportation fuel. There are other methods of making ethanol, and scientists have to consider many factors before deciding on the best method to produce it, including availability and cost of raw materials, energy requirements, rate of reaction, percentage yield, and atom economy of the reaction.

### The learning journey

Previously you will have learnt in SC9 *Calculations Involving Masses*:

- about the law of conservation of mass
- to calculate the maximum mass of a product formed from the mass of a reactant
- to calculate the concentration of a solution in  $\text{g dm}^{-3}$
- **H** to calculate the number of moles of a substance in a given mass of it and vice versa.

In this unit you will learn:

- reasons why the actual yield of a reaction is less than the theoretical yield
- to calculate the percentage yield of a reaction
- what is meant by the atom economy of a reaction and how to calculate it
- how to carry out an acid–alkali titration
- **H** how to calculate an unknown concentration or volume of a solution using titration
- **H** about the factors that are considered when selecting a manufacturing method
- **H** how to interconvert between  $\text{g dm}^{-3}$  and  $\text{mol dm}^{-3}$ .

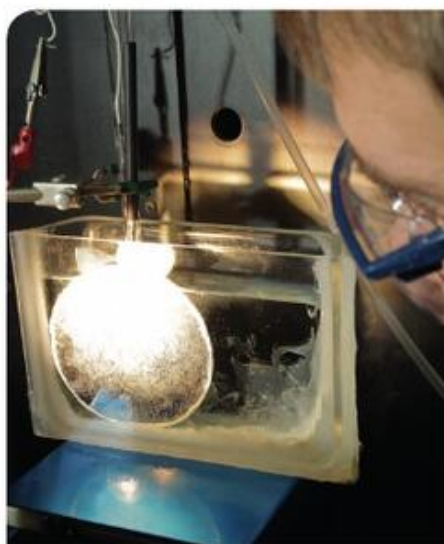


# SC14a Yields

Specification reference: C5.11C; C5.12C

## Progression questions

- What is meant by the terms theoretical yield and actual yield of a reaction?
- How do you calculate the percentage yield of a reaction?
- What are some reasons for the actual yield being less than the theoretical yield?



**A** Hydrogen is a potential fuel of the future but is very expensive to produce. Scientists have been working on using sunlight to split water into oxygen and hydrogen. A big advance was made in 2015 when scientists increased the **yield** of hydrogen by an order of magnitude (10 times) by using a substance called gallium phosphide.

If you split 36 g of water into oxygen and hydrogen, you should get 4 g of hydrogen. 4 g is the **theoretical yield** of hydrogen as it is the maximum mass of product that can be formed from the reactant. The theoretical yield is calculated from the balanced equation. In practice, you often do not get this much. The amount of product obtained when you carry out an experiment is known as the **actual yield**.

## Worked example W1

Calculate the theoretical yield of hydrogen from 36 g of water.

	$2\text{H}_2\text{O}$	$\rightarrow$	$2\text{H}_2$	$+ \text{O}_2$
Calculate relative formula masses and multiply by the balancing number	$2 \times (1 + 1 + 16) = 36$		$2 \times (1 + 1) = 4$	
Add the unit symbols	36 g		4 g	

So the theoretical yield of hydrogen is 4 g.

The **percentage yield** compares the actual yield and the theoretical yield:

$$\text{percentage yield} = \frac{\text{actual yield}}{\text{theoretical yield}} \times 100$$

## Worked example W2

In the reaction above, the actual yield was 3.3 g of hydrogen. Calculate the percentage yield.

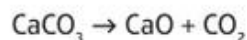
$$\text{percentage yield} = \frac{3.3}{4} \times 100 = 82.5\%$$

- 7<sup>th</sup>** 1 Calculate the percentage yield for a reaction where the actual yield is 3.2 g and the theoretical yield is 4.0 g.

## Did you know?

Scientists are trying to produce biodiesel by extracting oil from algae. The process is not very efficient at the moment, with only about 15% of the substances supplied to the algae being converted into useful oils.

- 2** When limestone is heated, the calcium carbonate decomposes to form calcium oxide:



- a** Calculate the theoretical yield of calcium oxide that could be made from 125 tonnes of calcium carbonate.



- b** The actual yield of calcium oxide is 45.5 tonnes. Calculate the percentage yield.



## Why is the yield less than expected?

When manufacturing substances, the theoretical yield is calculated assuming that all the reactants are turned into products, and that the products are successfully separated from the reaction mixture. There are three main reasons why reactions do not give 100% yields:

- the reaction may be incomplete so not all of the reactants are used up – possibly because the reaction has not been left for long enough, or the reaction may reach equilibrium
- some of the product is lost – for example, when a liquid is transferred from one container to another, some of it will be left behind on the walls of the first container
- there may be other unwanted **side reactions** taking place – for example, some of the reactants may react to make a different product. These side reactions compete with the main reaction.

The higher the percentage yield of a reaction, the more useful the reaction is. A high yield means that fewer raw materials are used to make the same amount of product, so there is less waste and more profit can be made.



**C** Calcium oxide is produced by heating limestone in a lime kiln. Modern lime kilns achieve a much greater percentage yield than old kilns, such as this Roman kiln.

- 3** Give a reason that explains why it is desirable to have a high percentage yield in a reaction.
- 4** One of the steps in the production of sulfuric acid is:
- $$2\text{SO}_2 + \text{O}_2 \rightarrow 2\text{SO}_3$$
- a** Calculate the theoretical yield of sulfur trioxide that could be obtained from 256 tonnes of sulfur dioxide.
- b** The actual yield of sulfur trioxide is 202 tonnes. Calculate the percentage yield.

### Exam-style question

In an experiment to make potassium chloride, the yield is 2.5 g. The theoretical yield of potassium chloride for this experiment is 4.0 g. Calculate the percentage yield of potassium chloride. (2 marks)



**B** When you bake a cake, some of the ingredients may get left behind on the scales, in the mixing bowl or in the cake tin. In a chemical reaction, some of the reactants and products may get left behind on the apparatus.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Carbon dioxide is formed when calcium carbonate reacts with hydrochloric acid. The theoretical yield for this reaction is 44 g. Calculate the percentage yield of carbon dioxide formed when the actual yield is 32 g.
- S2** Describe three reasons why the actual yield is less than the theoretical yield in a reaction.

### Extend

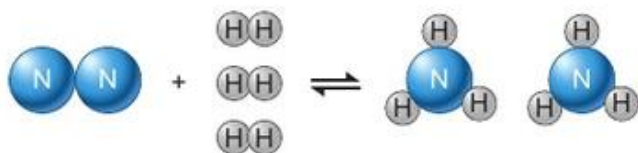
- E1** Iron is extracted from iron oxide by reducing it with carbon monoxide:
- $$\text{Fe}_2\text{O}_3 + 3\text{CO} \rightarrow 2\text{Fe} + 3\text{CO}_2$$
- a** Calculate the theoretical yield of iron that could be obtained from 320 tonnes of iron oxide.
- b** The actual yield of iron is 89.6 tonnes. Calculate the percentage yield.

# SC14b Atom economy

Specification reference: C5.13C; C5.14C; **H** C5.15C

## Progression questions

- What is meant by the atom economy of a reaction?
- How do you calculate the atom economy of a reaction?
- **H** How is data used to decide on the best way to manufacture a product?



**A** The atom economy for making ammonia is 100%.

**Atom economy** is a method of showing how efficiently a particular reaction makes use of the atoms in the reactants. Many reactions form more than one product and not all of the products are useful. The atom economy shows the percentage, by mass, of useful products.

## Did you know?

Ibuprofen was originally made in a six-step process with an atom economy of 40%. It is now made in a three-step process with an atom economy of 77%.

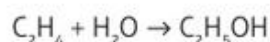


**B** Ibuprofen is a painkiller.

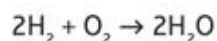
$M_r$  of ethanol is multiplied by 2 because the equation shows that one molecule of glucose produces two molecules of ethanol.

$$\text{atom economy} = \frac{\text{relative formula mass } (M_r) \text{ of the useful product}}{\text{sum of relative formula masses of all the reactants}} \times 100\%$$

There are two ways of making ethanol,  $\text{C}_2\text{H}_5\text{OH}$ . In one method ethene,  $\text{C}_2\text{H}_4$ , is reacted with steam. All of the atoms in the reactants are present in the product, so the atom economy is 100%.

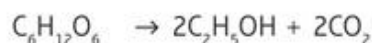


- 6<sup>th</sup>** 1 State what is meant by the atom economy of a reaction.
- 7<sup>th</sup>** 2 Show that the atom economy for making ethanol from ethene is 100%.
- 7<sup>th</sup>** 3 The following equation shows the reaction between hydrogen and oxygen.



What is the atom economy for making water in this way?

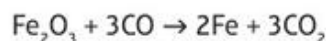
Ethanol is also produced by the fermentation of sugars, such as glucose. The atom economy of this method is much lower than making ethanol from ethene because carbon dioxide is formed as a **by-product**.



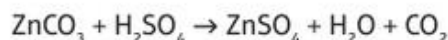
$$(M_r = 180) \quad (M_r = 46)$$

$$\text{atom economy for making ethanol} = \frac{2 \times 46}{180} \times 100 = 51.1\%$$

- 9<sup>th</sup>** 4 Calculate the atom economy for making iron in this reaction:



- 9<sup>th</sup>** 5 Zinc sulfate,  $\text{ZnSO}_4$ , can be made by reacting zinc carbonate with sulfuric acid:



Calculate the atom economy for making zinc sulfate in this way.



## H Reaction pathways

Chemists often have a choice of reaction pathway to produce a substance on a large scale.

A percentage yield calculation gives no indication of the amount of waste products. A reaction can have a high percentage yield but a low atom economy, meaning that waste by-products are formed. It is better to reduce the amount of waste produced than to have to treat it later.

One way to improve the atom economy of a reaction is to find uses for the by-products of the reaction.

Chemists must consider other factors when deciding on a reaction pathway, including energy consumption, rate of reaction, raw materials and the conditions needed to produce a high yield if the reaction reaches equilibrium.

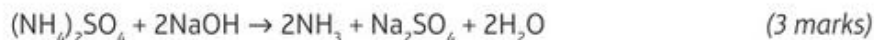
	Fermentation	Reaction of ethene with steam
raw materials	carbohydrates (e.g. from sugar cane, sugar beet, maize) – these are renewable	ethene obtained from crude oil – this is non-renewable
temperature	30–40 °C	300 °C
pressure	atmospheric pressure	high pressure (60–70 atm)
catalyst	enzymes in yeast	concentrated phosphoric acid
rate of reaction	slow	fast
purity	impure – must be fractionally distilled	pure

**D** comparison of two ways of making ethanol

- 7** **a** Compare the two methods of making ethanol in terms of raw materials, atom economy, energy consumption and the quality of ethanol produced.
- b** Brazil produces a lot of ethanol from sugar cane. Give three reasons that explain why Brazil produces ethanol from sugar cane instead of from ethene.

### Exam-style question

Calculate the atom economy for making ammonia from ammonium sulfate in the following reaction:



**C** In Edinburgh, Professor Martin Tangey and his team of scientists have developed a way of turning the waste products from whisky-making into butanol, for use as a biofuel.

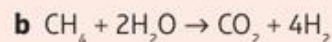
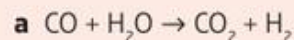
- 6** Explain, in terms of waste products, whether it is better for a reaction to have a high percentage yield and a low atom economy or a low percentage yield and a high atom economy.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Calculate the atom economy for producing hydrogen in the following reactions:



### Extend

**E1** **H** Explain how atom economy, percentage yield and the usefulness of waste products will affect the choice of reaction pathway for manufacturing a particular product.

# SC14c Concentrations

Specification reference: **H** C5.8C

## Progression questions

- **H** How do you calculate the concentration of a solution in  $\text{g dm}^{-3}$ ?
- **H** How do you calculate the concentration of a solution in  $\text{mol dm}^{-3}$ ?
- **H** How do you convert a concentration in  $\text{g dm}^{-3}$  into  $\text{mol dm}^{-3}$  and vice versa?

**H**



**A** A volumetric flask is used for making an accurate solution.

A solution with an accurate concentration is made up in a **volumetric flask**. These flasks are **calibrated** to measure one volume of solution accurately, for example,  $100.0\text{ cm}^3$ ,  $250.0\text{ cm}^3$  or  $1.00\text{ dm}^3$ . To make a solution, the solute is dissolved in distilled water and then made up to the graduation mark before stoppering and then shaking the flask.

It is important to measure the volume from the bottom of the meniscus and to look at it with your eyes on the same level as the graduation mark to get an accurate measurement and avoid a random error.

The concentration of a solution is the amount of solute dissolved in a stated volume of solution. The units of concentration are usually written as  $\text{g dm}^{-3}$ .  $1\text{ dm}^3$  is the same volume as 1 litre or  $1000\text{ cm}^3$ .



**B** Fill the flask so the bottom of the meniscus is on the graduation mark.

You can calculate the concentration of a solution in  $\text{g dm}^{-3}$  using this equation:

$$\text{concentration in } \text{g dm}^{-3} = \frac{\text{mass of solute in g}}{\text{volume of solution in } \text{dm}^3}$$

If the volume of the solution is given in  $\text{cm}^3$ , convert it to  $\text{dm}^3$  by dividing by 1000. You can use an equation triangle to help re-arrange this equation.

- 1 Calculate the concentration, in  $\text{g dm}^{-3}$ , of the solute in these solutions:



**a** 15 g of sodium chloride in  $3.0\text{ dm}^3$  of solution



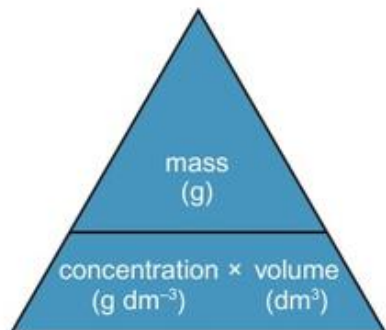
**b** 0.25 g of sodium hydroxide in  $100\text{ cm}^3$  of solution.



- 2 Calculate the mass of magnesium sulfate in  $50\text{ cm}^3$  of a solution of concentration  $10\text{ g dm}^{-3}$ .



- 3 Calculate the volume of solution, in  $\text{cm}^3$ , containing 0.5 g of zinc nitrate that will have a concentration of  $2.5\text{ g dm}^{-3}$ .



**C** equation triangle for working out concentration



## H Concentrations in mol dm<sup>-3</sup>

The concentration of a solution is often given as the number of moles of solute dissolved in 1 dm<sup>3</sup> of solution. The unit is written as mol dm<sup>-3</sup>.

You can calculate the concentration of a solution in mol dm<sup>-3</sup> using this equation:

$$\text{concentration in mol dm}^{-3} = \frac{\text{number of moles of solute}}{\text{volume of solution in dm}^3}$$

- 4 Calculate the concentration, in mol dm<sup>-3</sup>, of the solute in these solutions:



a 1.0 mol of potassium chloride in 4.0 dm<sup>3</sup> of solution



b 0.15 mol of copper sulfate in 250 cm<sup>3</sup> of solution.



- 5 Calculate the number of moles of sodium sulfate in 100 cm<sup>3</sup> of a solution of concentration 0.25 mol dm<sup>-3</sup>.



- 6 Calculate the volume of solution, in cm<sup>3</sup>, containing 0.02 mol of potassium hydroxide that will have a concentration of 0.5 mol dm<sup>-3</sup>.

The two types of concentrations are related by the equation:

$$\text{concentration in mol dm}^{-3} = \frac{\text{concentration in g dm}^{-3}}{\text{relative formula mass of solute}}$$

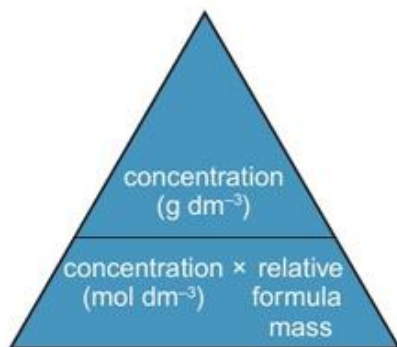
- 7 Calculate the concentration, in mol dm<sup>-3</sup>, of these solutions:



a sulfuric acid, H<sub>2</sub>SO<sub>4</sub>, of concentration 98 g dm<sup>-3</sup>



b hydrochloric acid, HCl, of concentration 1.825 g dm<sup>-3</sup>.



E equation triangle for converting concentrations

- 8 Calculate the concentration, in g dm<sup>-3</sup>, of these solutions:



a sodium hydroxide, NaOH, of concentration 0.5 mol dm<sup>-3</sup>



b sodium chloride, NaCl, of concentration 0.125 mol dm<sup>-3</sup>.

### Exam-style question

Calculate the concentration, in g dm<sup>-3</sup>, of a solution of nitric acid, HNO<sub>3</sub>, with a concentration of 0.4 mol dm<sup>-3</sup>. (2 marks)

### Did you know?

Some chemists celebrate 'Mole Day' between 6.02 am and 6.02 pm on 23 October each year.



### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1 A solution of sodium hydroxide, NaOH, has a concentration of 32 g dm<sup>-3</sup>. Calculate the concentration of the solution in mol dm<sup>-3</sup>.
- S2 A solution of sodium carbonate, Na<sub>2</sub>CO<sub>3</sub>, has a concentration of 0.1 mol dm<sup>-3</sup>. Calculate the concentration of the solution in g dm<sup>-3</sup>.

### Extend

- E1 A student needs to make up a solution of potassium carbonate, K<sub>2</sub>CO<sub>3</sub>, of concentration 0.05 mol dm<sup>-3</sup>. Calculate the mass of potassium carbonate needed to make up 250 cm<sup>3</sup> of this solution.
- E2 A solution of hydrochloric acid has a concentration of 0.1 mol dm<sup>-3</sup>. Calculate the number of moles of HCl in a 25.0 cm<sup>3</sup> portion of this solution.

# SC14d Titrations and calculations

Specification reference: C5.9C; **H** C5.10C

## Progression questions

- How do you carry out an acid–alkali titration?
- **H** How do you calculate the number of moles of solute in a given volume of solution?
- **H** How do you calculate the concentration of a solution using the results of an acid–alkali titration?

- 6<sup>th</sup>** **1** 25.0 cm<sup>3</sup> of sodium hydroxide solution was placed in a conical flask with a few drops of methyl orange. It was titrated with hydrochloric acid. State the indicator colour change at the end-point.

Acid–alkali titrations are used to find the exact volume of an acid that neutralises a specified volume of an alkali or vice versa. Acids and alkalis are colourless so we use indicators. The apparatus is shown in *SC14d Core practical*.

**H**

The concentration of one of the solutions in a titration can be calculated if the concentration of the other solution is known.

$$\text{concentration in mol dm}^{-3} = \frac{\text{number of moles of solute}}{\text{volume of solution in dm}^3}$$

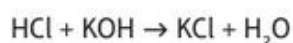
The **mole ratio** is the ratio in moles of the substances in a balanced equation.

## Did you know?

Titrations have many uses including: testing water samples, and analysing foodstuffs and drugs.

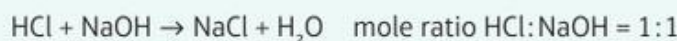
- 9<sup>th</sup>** **2** Sulfuric acid, H<sub>2</sub>SO<sub>4</sub>, reacts with a solution of potassium hydroxide, KOH. Write the balanced equation for this reaction.

- 9<sup>th</sup>** **3** 25.0 cm<sup>3</sup> of potassium hydroxide solution was neutralised by 35.0 cm<sup>3</sup> of 0.750 mol dm<sup>-3</sup> dilute hydrochloric acid. Calculate the concentration, in mol dm<sup>-3</sup>, of the potassium hydroxide solution used.



## Worked example W1

25.0 cm<sup>3</sup> of sodium hydroxide solution was titrated against 0.100 mol dm<sup>-3</sup> hydrochloric acid. An average of 20.0 cm<sup>3</sup> of the acid neutralised the alkali. Calculate the concentration of the sodium hydroxide solution.



Step 1: calculate the number of moles of the solution of known volume and concentration.

$$\begin{aligned} \text{number of moles of HCl} &= \text{concentration of HCl} \times \text{volume used} \\ & \quad (\text{mol dm}^{-3}) \quad (\text{dm}^3) \\ &= 0.100 \times \frac{20.0}{1000} = 0.00200 \text{ mol} \end{aligned}$$

Step 2: use the balanced equation to work out the number of moles of alkali that reacted.

1 mol of HCl reacts with 1 mol of NaOH, so 0.00200 mol of HCl reacts with 0.00200 mol of NaOH.

Step 3: calculate the concentration of the sodium hydroxide solution.

$$\text{concentration of NaOH in mol dm}^{-3} = \frac{\text{number of moles of NaOH}}{\text{volume of NaOH solution in dm}^3}$$

$$\text{volume of NaOH solution} = \frac{25.0}{1000} = 0.0250 \text{ dm}^3$$

$$\text{concentration of NaOH} = \frac{0.00200}{0.0250} = 0.0800 \text{ mol dm}^{-3}$$

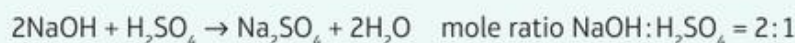


**H**

These types of calculations can also be used to calculate the volume of one solution that will react with a given volume of another. You need to know the concentrations of both solutions and the balanced equation.

**Worked example W2**

25.0 cm<sup>3</sup> of 0.200 mol dm<sup>-3</sup> sodium hydroxide solution was neutralised by 0.150 mol dm<sup>-3</sup> sulfuric acid. Calculate the volume of sulfuric acid needed for the neutralisation.



Step 1: calculate the number of moles of the solution of known volume and concentration.

$$\begin{aligned} \text{number of moles of NaOH} &= \text{concentration of NaOH} \times \text{volume used} \\ &= 0.200 \times \frac{25.0}{1000} = 0.00500 \text{ mol} \end{aligned}$$

Step 2: use the balanced equation and mole ratio to work out the number of moles of acid that reacted.

2 mol of NaOH reacts with 1 mol of H<sub>2</sub>SO<sub>4</sub>,

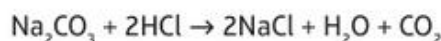
so 0.00500 mol of NaOH reacts with  $\frac{0.00500}{2} = 0.00250$  mol of H<sub>2</sub>SO<sub>4</sub>

Step 3: calculate the volume of sulfuric acid needed.

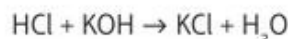
$$\begin{aligned} \text{volume in dm}^3 &= \frac{\text{number of moles of acid}}{\text{concentration in mol dm}^{-3}} \\ &= \frac{0.00250}{0.150} = 0.0167 \text{ dm}^3 \text{ or } 16.7 \text{ cm}^3 \end{aligned}$$

Round the final answer to the same number of significant figures as used in the question.

- 6** 10.0 cm<sup>3</sup> of 0.095 mol dm<sup>-3</sup> sodium carbonate solution was neutralised by 0.15 mol dm<sup>-3</sup> hydrochloric acid. Calculate the volume of hydrochloric acid, in cm<sup>3</sup>, needed.

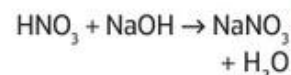

**Exam-style question**

25.0 cm<sup>3</sup> of potassium hydroxide solution reacted with 23.3 cm<sup>3</sup> of 0.100 mol dm<sup>-3</sup> hydrochloric acid. Calculate the concentration of this potassium hydroxide solution in mol dm<sup>-3</sup>.

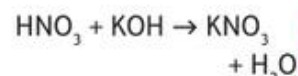


(3 marks)

- 4** 10.0 cm<sup>3</sup> of 0.140 mol dm<sup>-3</sup> sodium hydroxide solution was neutralised by 12.3 cm<sup>3</sup> of nitric acid. Calculate the concentration, in mol dm<sup>-3</sup>, of the nitric acid.



- 5** 25.0 cm<sup>3</sup> of 0.275 mol dm<sup>-3</sup> potassium hydroxide solution was neutralised by 0.166 mol dm<sup>-3</sup> nitric acid. Calculate the volume of nitric acid, in cm<sup>3</sup>, needed to neutralise the potassium hydroxide solution.


**Checkpoint**

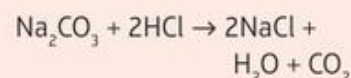
How confidently can you answer the Progression questions?

**Strengthen**

- S1** Describe how to carry out a titration to find the exact volume of hydrochloric acid needed to neutralise 25.0 cm<sup>3</sup> of a sodium hydroxide solution.

**Extend**

- E1** **H** 25.0 cm<sup>3</sup> of sodium carbonate solution was neutralised by 22.6 cm<sup>3</sup> of 0.100 mol dm<sup>-3</sup> hydrochloric acid. Calculate the concentration of the sodium carbonate solution in g dm<sup>-3</sup>.



# SC14d Core practical – Acid–alkali titration

Specification reference: C5.9C

## Aim

Carry out an accurate acid–alkali titration, using a burette, a pipette and a suitable indicator.



A a titration experiment



B The initial volume of solution in the burette is 0.20 cm<sup>3</sup> and the final burette reading is 22.20 cm<sup>3</sup>.



C Methyl orange indicator is yellow in alkalis, peachy-orange in neutral solutions and pink in acids.

Titration is used to find the exact volume of one solution that reacts with a fixed volume of another solution. The fixed volume of solution, often 25.0 cm<sup>3</sup>, is measured in a pipette and the other solution is contained in a burette as these pieces of apparatus give accurate measurements. A burette has a fine scale, which gives it a good resolution. It can be read to the nearest half division, that is to  $\pm 0.05$  cm<sup>3</sup>. Remember that the measuring instruments are calibrated for readings taken from the *bottom* of the meniscus.

The result of an acid–alkali titration can be used to prepare a soluble salt or to calculate the concentration of a solution.

## Your task

You will carry out a titration to find the exact volume of hydrochloric acid needed to neutralise 25.0 cm<sup>3</sup> of a solution of sodium hydroxide.

## Method

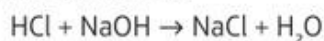
Wear eye protection. Avoid skin contact with the liquids.

- A** Rinse a burette with hydrochloric acid, then fill the burette with the acid, making sure the jet below the tap is also full.
- B** Record the initial volume of acid in the burette.
- C** Rinse a pipette with sodium hydroxide solution, then fill the pipette to the 25.0 cm<sup>3</sup> mark and empty the solution into a conical flask.
- D** Add a few drops of methyl orange indicator to the flask and place the flask on a white tile under the burette.
- E** Add the acid to the sodium hydroxide solution while swirling the flask.
- F** When the indicator starts to change colour, rinse the tip of the burette and the sides of the flask with a small amount of distilled water from a wash bottle to ensure that all the acid is in the mixture, then add the acid drop by drop until the end-point is reached.
- G** Record the final volume of acid in the burette.
- H** Repeat the experiment, apart from the initial rinsing of the burette and pipette, until concordant results are obtained.



## Exam-style questions

- 1 State a safety precaution to take in this practical and the reason for it. (1 mark)
- 2 Give a reason why the burette and pipette should be rinsed before they are filled. (1 mark)
- 3 State what you should do after filling the burette with acid and before taking the initial reading. (1 mark)
- 4 Give a reason for standing the conical flask on a white tile. (1 mark)
- 5 Suggest a reason why universal indicator is not suitable for a titration. (1 mark)
- 6 State what is meant by the 'end-point' of the titration. (1 mark)
- 7 State the colour change that would be seen at the end-point in this titration using methyl orange indicator. (2 marks)
- 8 State what is meant by 'concordant results'. (1 mark)
- 9 25.0 cm<sup>3</sup> of sodium hydroxide solution was titrated with dilute hydrochloric acid:



The following results were obtained.

	Titration 1	Titration 2	Titration 3
final burette reading (cm <sup>3</sup> )	26.00	26.30	27.20
initial burette reading (cm <sup>3</sup> )	0.00	1.20	1.90
volume of acid used (cm <sup>3</sup> )	26.00	25.10	

- a Calculate the volume of acid used in titration 3. (1 mark)
  - b Calculate the volume of acid that should be used to react with 25.0 cm<sup>3</sup> of the sodium hydroxide solution to produce the salt sodium chloride. Give a reason for your answer. (2 marks)
  - c Describe how you would use the volume calculated in **b** to obtain pure, dry crystals of sodium chloride. (3 marks)
  - d **H** The sodium hydroxide solution had a concentration of 0.100 mol dm<sup>-3</sup>. Calculate the concentration, in mol dm<sup>-3</sup>, of the hydrochloric acid used in this experiment. (3 marks)
- 10 **H** Give a reason for using a burette, rather than a measuring cylinder, for measuring the volume of hydrochloric acid.
  - 11 **H** A student carried out a titration experiment following the method given, but measured the volumes of the solutions in the burette and the pipette from the top of the meniscus instead of the bottom. Explain how this error will affect the volumes measured, if at all. (3 marks)

# SC14e Molar volume of gases

Specification reference: **H** C5.16C; **H** C5.17C; **H** C5.18C

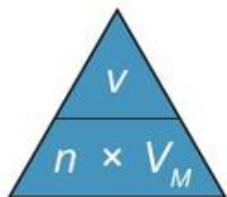
## Progression questions

- **H** What is Avogadro's law?
- **H** What is the molar volume of a gas?
- **H** How do you calculate the volume of a gas, and the mass of a solid, involved in a chemical reaction?

**H**



**A** In 1914, one way to manufacture hydrogen was the reaction between silicon and hot, concentrated sodium hydroxide solution.

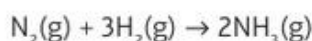


**B**  $n$  = amount in mol,  $v$  = volume of gas,  
 $V_M$  = molar volume

## Avogadro's law

The volume occupied by a sample of a gas depends on the temperature, pressure and number of particles of gas. The volume of a gas does *not* depend on its relative formula mass,  $M_r$ . Remember that the mean distance between gas particles is very large compared to the size of the particles.

**Avogadro's law** states that, if the temperature and pressure are the same, equal volumes of different gases contain an equal number of molecules. This lets you calculate the volumes of gases involved in a reaction. For example, nitrogen and hydrogen react together to form ammonia:



The ratio of nitrogen to ammonia is 1:2. So 100 cm<sup>3</sup> of nitrogen will produce 200 cm<sup>3</sup> of ammonia, at the same temperature and pressure, if it reacts completely with hydrogen.



- 1** State two assumptions needed when using Avogadro's law in calculations.



- 2** Calculate the volume of hydrogen that will react completely with 100 cm<sup>3</sup> of nitrogen.

## Molar gas volume

The **molar gas volume** is the volume occupied by one mole of molecules of any gas. It is 24 dm<sup>3</sup> or 24 000 cm<sup>3</sup> at room temperature and pressure (rtp):

$$\text{volume of gas} = \text{amount of gas (mol)} \times \text{molar volume}$$

For example, 0.5 mol of carbon dioxide occupies  $0.5 \times 24 = 12 \text{ dm}^3$  at rtp.



- 3** Calculate the volume, in cm<sup>3</sup>, occupied by 0.1 mol of oxygen at rtp.

If you know the volume of a gas at rtp, you can calculate its amount:

$$\text{amount of gas (mol)} = \frac{\text{volume of gas}}{\text{molar volume}}$$

For example, 96 cm<sup>3</sup> of hydrogen contains  $\frac{96}{24\,000} = 4.0 \times 10^{-3}$  mol of hydrogen molecules.

You can use calculations like these to calculate the masses of solids and volumes of gases in chemical reactions.



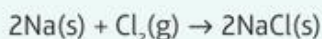
- 4** Calculate the amount of chlorine molecules, in mol, in 16 dm<sup>3</sup> of chlorine. Give your answer to 2 significant figures.



## H

## Worked examples

- 1 Sodium reacts with chlorine to form sodium chloride:



- a Calculate the minimum volume at rtp, in  $\text{dm}^3$ , of chlorine needed to react completely with 0.92 g of sodium.

$$\text{amount of sodium} = \frac{0.92}{23} = 0.040 \text{ mol}$$

From the balanced equation, 2 mol of Na reacts with 1 mol of  $\text{Cl}_2$

$$\text{So } 0.040 \text{ mol of Na reacts with } \frac{0.040}{2} = 0.020 \text{ mol of } \text{Cl}_2$$

$$\text{volume of } \text{Cl}_2 = 0.020 \times 24 = 0.48 \text{ dm}^3$$

- b Calculate the maximum mass of sodium chloride that can be made from 960  $\text{cm}^3$  of chlorine at rtp and an excess of sodium.

$$\text{amount of chlorine} = \frac{960}{24\,000} = 0.040 \text{ mol}$$

From the balanced equation, 1 mol of  $\text{Cl}_2$  produces 2 mol of NaCl

$$\text{So } 0.040 \text{ mol of } \text{Cl}_2 \text{ produces } 2 \times 0.040 = 0.080 \text{ mol of NaCl}$$

$$\text{mass of NaCl} = 0.080 \times 58.5 = 4.68 \text{ g (4.7 g to 2 significant figures)}$$

- 2 Copper carbonate decomposes when heated:



Calculate the minimum mass of copper carbonate needed to produce 1.44  $\text{dm}^3$  of carbon dioxide at rtp.

$$\text{amount of carbon dioxide} = \frac{1.44}{24} = 0.060 \text{ mol}$$

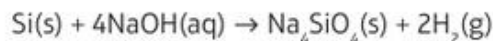
From the balanced equation, 1 mol of  $\text{CO}_2$  is made from 1 mol of  $\text{CuCO}_3$

$$\text{So } 0.060 \text{ mol of } \text{CO}_2 \text{ needs } 1 \times 0.060 = 0.060 \text{ mol of } \text{CuCO}_3$$

$$\text{mass of } \text{CuCO}_3 = 0.060 \times 123.5 = 7.41 \text{ g (7.4 g to 2 significant figures)}$$

## Exam-style question

Silicon reacts with hot, concentrated sodium hydroxide solution to produce sodium silicate and hydrogen:

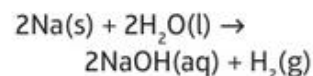


Calculate the minimum mass of silicon needed to produce 1000  $\text{dm}^3$  of hydrogen at rtp. (2 marks)



C Hot sodium burns in chlorine with a yellow flame, producing sodium chloride.

- 5 Sodium reacts with water to produce sodium hydroxide and hydrogen:



- a Calculate the maximum volume at rtp, in  $\text{dm}^3$ , of hydrogen that can be produced from 4.6 g of sodium.



- b Calculate the minimum mass of water needed to produce 0.36  $\text{dm}^3$  of hydrogen at rtp.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1 How do you calculate the volumes of gases involved in chemical reactions?

## Extend

- E1 Flying in the 1930s, the *Hindenburg* was the largest ever airship. Calculate the mass of zinc that would be needed, when reacted with excess sulfuric acid, to produce the 200 000  $\text{m}^3$  of hydrogen it contained.

# SC15a Fertilisers and the Haber process

Specification reference: C5.19C; C5.22C; C5.23C; C5.24C

## Progression questions

- What are fertilisers?
- What are the similarities and differences between making a fertiliser in a laboratory and in a factory?
- How is the Haber process used in the manufacture of ammonium nitrate?



**A** The corn cob on the left has been grown using a nitrogen-rich fertiliser, while the one on the right has not.

Plants absorb water and mineral ions through their root hair cells. This gradually reduces the concentration of mineral ions in the soil. Plants may then not grow properly and may suffer from deficiency diseases, such as stunted shoots and discoloured leaves.

**Fertilisers** replace the mineral ions needed by plants and so promote plant growth. Nitrogen, phosphorus and potassium (N, P and K) are important elements, but fertilisers must provide them as soluble compounds because root hair cells only absorb mineral ions that are dissolved in water.



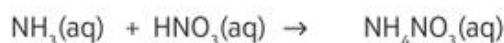
- 1** Explain why potassium nitrate,  $\text{KNO}_3$ , and ammonium phosphate,  $(\text{NH}_4)_3\text{PO}_4$ , are suitable compounds to use in fertilisers.

## Ammonium nitrate

**Nitrogenous** fertilisers are a source of soluble nitrogen compounds.

Ammonium nitrate is one such nitrogen-rich fertiliser. Ammonium nitrate is a salt manufactured by reacting ammonia solution with dilute nitric acid:

ammonia + nitric acid  $\rightarrow$  ammonium nitrate



Ammonia is manufactured by the Haber process, in which nitrogen and hydrogen react together in a **reversible reaction** (see SC12a *Dynamic equilibrium*).

Nitric acid is manufactured from ammonia in stages. Overall, the reaction may be represented as:

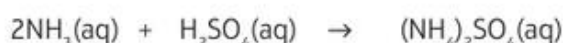
ammonia + oxygen  $\rightarrow$  nitric acid + water



## Ammonium sulfate

Ammonium sulfate is also a nitrogenous fertiliser. It is made in the laboratory by reacting ammonia solution with dilute sulfuric acid:

ammonia + sulfuric acid  $\rightarrow$  ammonium sulfate



Preparing ammonium sulfate in the laboratory is very different from its industrial production. Table B shows some differences.

## Did you know?

The annual world production of ammonia by the Haber process is around 175 million tonnes (about 24 kg per person per year). About 85% of this is used to make nitrogenous fertilisers.

- 2** State and explain the type of reaction involved in making:



**a** ammonium nitrate



**b** nitric acid.



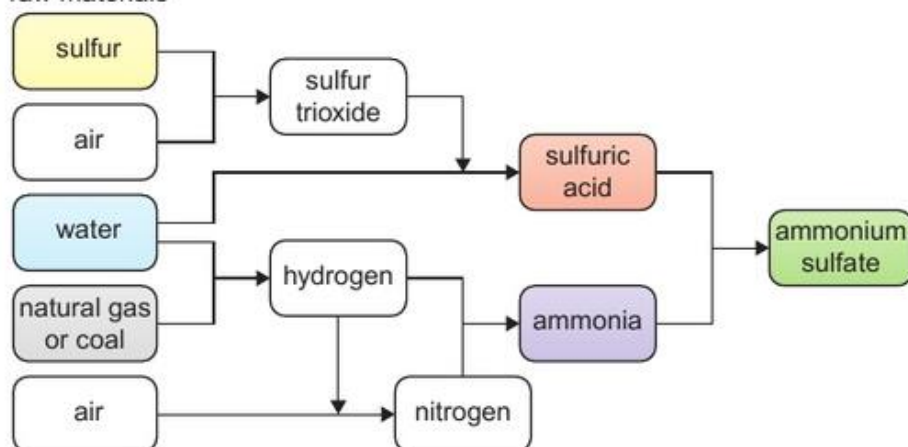


	Laboratory preparation	Industrial production
scale of production	small scale	large scale
starting materials	ammonia solution and dilute sulfuric acid	raw materials for making ammonia and sulfuric acid
stages	titration (see <i>SC8 Acids and Alkalis</i> ), then crystallisation	several stages
type of process	batch	continuous

**B** some features of making ammonium sulfate in the laboratory or industrially

For the laboratory preparation of ammonium sulfate, ammonia solution and dilute sulfuric acid are bought from chemical manufacturers. For industrial production, these substances are often made on-site from their raw materials (see diagram C).

raw materials



**C** flow chart for the industrial production of ammonium sulfate

The laboratory preparation of ammonium sulfate is a **batch process**. A small amount is made, the apparatus is cleaned, and then another small amount is made. Industrial preparation is a **continuous process**, in which reactants are constantly fed into the reactors and products are removed. The process is stopped only rarely to maintain and clean the equipment. Batch processes are difficult to automate. However, continuous processes are usually automated, so they need fewer people to make a given amount of product.

- 5** The rate at which each reactant is fed into the reactor in a continuous process must be carefully controlled. Suggest reasons that explain this.
- 6** Describe two advantages of a continuous process compared to a batch process.

- 3** Use diagram C to name the raw materials needed to make sulfuric acid and ammonia in industry.
- 4** One of the stages in the manufacture of sulfuric acid is a reversible reaction. Sulfur dioxide,  $\text{SO}_2$ , reacts with oxygen from the air to produce sulfur trioxide,  $\text{SO}_3$ . Write a balanced equation, including state symbols, for this reaction.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** How are nitrogenous fertilisers made in the laboratory?
- S2** How are nitrogenous fertilisers made on an industrial scale?

### Extend

- E1** Compare the advantages and disadvantages of ammonia and ammonium sulfate as nitrogenous fertilisers.

### Exam-style question

Suggest reasons that explain why farmers may apply 'NPK' fertilisers to their fields. (2 marks)

# SC15b Factors affecting equilibrium

Specification reference: **H** C5.15C; **H** C5.20C; **H** C5.21C

## Progression questions

- **H** How is the time taken to reach equilibrium affected by changes in conditions?
- **H** How are conditions chosen for industrial chemical reactions?
- **H** How are reaction pathways chosen for industrial processes?

**H**



**A** Nitric acid is made using nitrogen monoxide, produced from nitrogen and oxygen in an endothermic reaction. Nitrogen monoxide is made during thunderstorms because lightning causes the very high temperatures needed to move the equilibrium position in the direction of the endothermic reaction.



- 1** Compare the effect of catalysts on equilibria with the effects of changing temperature, pressure or concentration.



- 2** Why might an industrial reaction *not* be allowed to reach equilibrium?

Reversible reactions may reach equilibrium (see SC12a *Dynamic equilibrium*). This only happens when they are in a **closed system**, such as a stoppered flask, in which no substances can enter or leave.

In a **dynamic equilibrium**:

- the forward and backward reactions still happen, and at the same rate
- the concentrations of all reacting substances do not change.

The position of a dynamic equilibrium, and how quickly equilibrium is reached, are affected by changes in conditions (see table B).

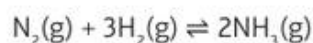
Change in conditions	Position of equilibrium	Time taken to reach equilibrium
temperature increased	moves in the direction of the endothermic reaction	decreases
pressure increased in a reaction involving gases	moves towards the side of the balanced equation with the fewer molecules of reacting gas	decreases
concentration of a reacting substance increased	moves away from the reacting substance in the balanced equation	decreases
catalyst added	no change	decreases

**B** effects of reaction conditions on dynamic equilibria

## Equilibria and industrial processes

Chemical manufacturers must make a profit. They choose reaction conditions to produce an acceptable product **yield** in an acceptable time. This may mean that reactions are not allowed to reach equilibrium – it is not profitable to achieve equilibrium if it takes years. In addition, choosing conditions that give a high rate of reaction is not worthwhile if they are too expensive.

In the Haber process, nitrogen and hydrogen react together to produce ammonia:



There are fewer molecules of reacting gas on the right-hand side of the equation. As the pressure is increased, the position of equilibrium moves to the right.



**H**

High pressures increase the equilibrium yield of ammonia *and* increase the rate of reaction. The pressure chosen, 200 atmospheres, is a compromise because very high pressures are too expensive to maintain.

The forward reaction is exothermic, so the backward reaction is endothermic. As the temperature is increased, the position of equilibrium moves to the left (in the direction of the endothermic reaction). Low temperatures increase the equilibrium yield of ammonia but reduce the rate of reaction. The temperature chosen, 450 °C, is a compromise that gives an acceptable yield of ammonia in an acceptable time.




An iron catalyst is used because, although it does not change the position of equilibrium, it *does* increase the rate of reaction.

### Choosing reaction pathways

There is usually more than one way to make a product. A particular **reaction pathway** is chosen by taking into account factors such as:

- availability and cost of raw materials and energy supplies
- the rate of reaction and the equilibrium position
- atom economy, yield and usefulness of **by-products** (see SC14b Atom economy).

In the past, nitric acid,  $\text{HNO}_3$ , was made using the Birkeland–Eyde process. This imitated the effect of lightning (photo A). A high-voltage electric arc was created between two electrodes. Nitrogen and oxygen in the air reacted to form nitrogen monoxide, NO. This further reacted with oxygen to form nitrogen dioxide,  $\text{NO}_2$ , which was then dissolved in water to form nitric acid. The process used huge amounts of electricity and the **yield** of nitrogen monoxide was only around 4%. Today, nitric acid is made from ammonia using the Ostwald process (diagram C).

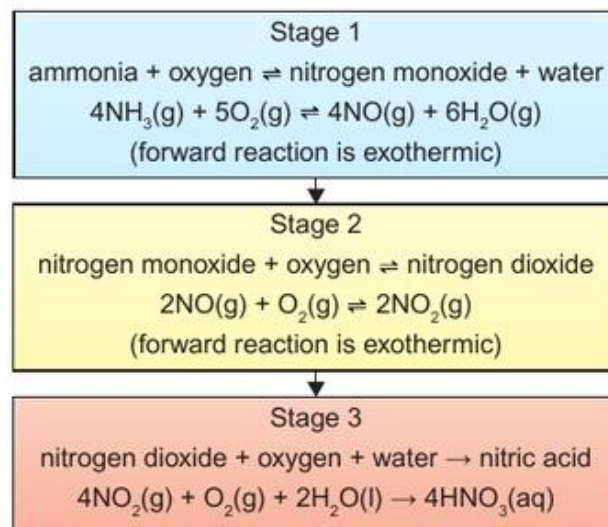
-  **4** Compare stages 2 and 3 of the Ostwald process with the Birkeland–Eyde process for making nitric acid.
-  **5** Compare and contrast stage 1 of the Ostwald process with the Birkeland–Eyde process.
-  **6** Explain the reaction conditions chosen for stage 1 of the Ostwald process.

### Exam-style question

Stage 2 of the Ostwald process happens at about 50 °C (see diagram C). State and explain what this shows about the rate of the reaction at stage 2. (2 marks)



- 3** Explain why the Haber process does not use very high pressures or low temperatures.



**C** Flow chart for the industrial production of nitric acid using the Ostwald process. Stage 1 is carried out at around 220 °C and 4 atmospheres of pressure with hot platinum as the catalyst. Under these conditions, the yield of nitrogen monoxide is around 95%. The reactions are so exothermic that the gases must be cooled.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** How does the time taken for a reversible reaction to reach equilibrium depend upon the reaction conditions?
- S2** How are the reaction conditions chosen for industrial processes?

### Extend

- E1** Evaluate the use of high temperatures and pressures for industrial reactions.

# SC16a Chemical cells and fuel cells

Specification reference: C5.25C; C5.26C; C5.27C

## Progression questions

- Why do batteries go 'flat'?
- What happens in a hydrogen–oxygen fuel cell?
- What are the strengths and weaknesses of fuel cells?



**A** This lithium ion battery pack is being moved into position, ready to fit into an electric car.



- 1** Suggest a reason that explains why the Daniell cell is described as a 'wet cell'.

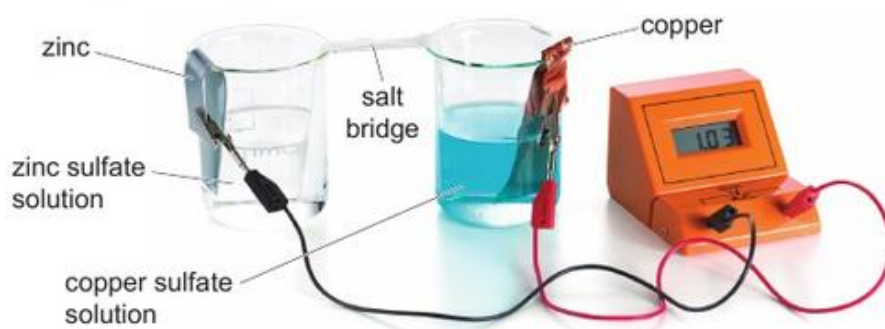


- 2** Explain what happens to the mass of each metal when a Daniell cell is in use.

The everyday batteries used in mobile phones and torches are **chemical cells**. A simple chemical cell has these components:

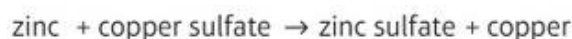
- two different metals, each dipped into a solution of one of their salts
- a 'salt bridge' to allow dissolved ions to pass from one solution to the other.

A voltage (potential difference) is produced between the two metals. In general, the further apart in the reactivity series the two metals are, the greater the voltage. A current flows if the cell is connected to an external circuit. Photo B shows a simple chemical cell, invented in 1836 by John Daniell (1790–1845). The salt bridge is filter paper soaked with concentrated potassium nitrate solution.



**B** This Daniell cell is connected to a voltmeter, which measures its voltage.

The overall reaction that happens in a Daniell cell is:



This is an exothermic reaction, but the energy is transferred mainly by electricity rather than by heating. When one of the reactants is used up, the reaction stops and a voltage is no longer produced. This is why household batteries go 'flat', and why electric cars need recharging after a journey.

Most modern batteries are 'dry cells' containing various reactants. Their solutions are mixed with a powder to make a paste.

## Fuel cells

Chemical cells store all their reactants, but **fuel cells** are supplied with fuel and oxygen from outside. No burning takes place inside fuel cells, just like in chemical cells. Fuel cells do not go 'flat' – they produce a voltage for as long as reactants are supplied. Hydrogen–oxygen fuel cells use hydrogen and oxygen, and water is the only product. Diagram C shows what happens.



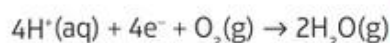
- 3** Write a balanced equation for the overall reaction in a hydrogen–oxygen fuel cell.

**H**

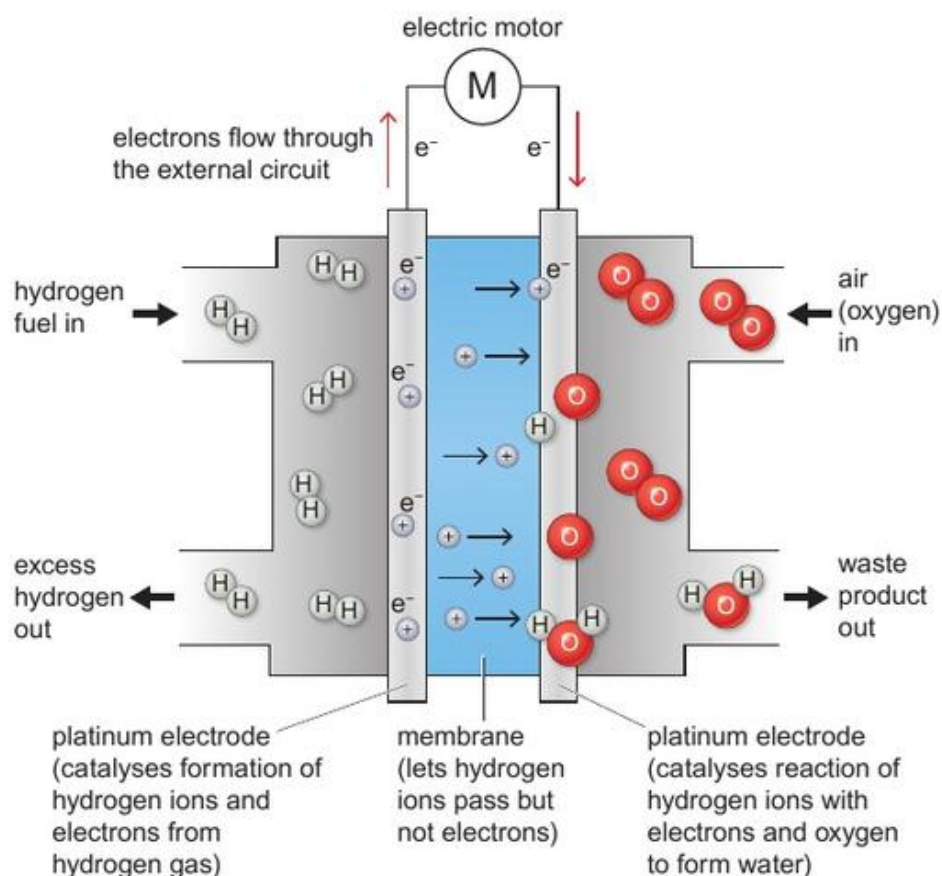
At the left-hand electrode, hydrogen atoms lose electrons and form hydrogen ions:



Electrons flow through the external circuit to the positive electrode. Hydrogen ions pass through a membrane to the right-hand electrode, where they gain electrons and react with oxygen:

**4**

- Explain whether hydrogen is oxidised or reduced at the left-hand electrode.



**C** a hydrogen–oxygen fuel cell with an electric motor in the external circuit

A hydrogen–oxygen fuel cell and electric motor are much quieter and need less maintenance than a petrol or diesel engine, but the hydrogen still needs to be stored in a tank.

Hydrogen–oxygen fuel cells do not produce carbon dioxide (a **greenhouse gas**). However, most hydrogen is manufactured by the reaction of steam with coal or natural gas. These processes do release carbon dioxide as a **by-product**.

- 5** Explain why a hydrogen–oxygen fuel cell car may cause the release of greenhouse gases.
- 6** In 2015, there were several thousand filling stations in the UK supplying petrol and diesel, but only four supplying hydrogen to the public. Explain why this limits the appeal of hydrogen–oxygen fuel cell cars, even though their range is similar to that of conventional cars.

### Exam-style question

Some fuel cells use ethanol,  $\text{C}_2\text{H}_5\text{OH}$ . When in use, all the carbon atoms are converted to carbon dioxide. Write a balanced equation for the overall reaction in an ethanol–oxygen fuel cell. (2 marks)

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** How are chemical reactions used to produce electricity?
- S2** What are the strengths and weaknesses of using a hydrogen–oxygen fuel cell to power a car's electric motor?

### Extend

- E1** Suggest reasons that explain why hydrogen–oxygen fuel cells and solar cells are used to produce electricity for the International Space Station.

## Making a fertiliser

Ammonium sulfate is useful as a fertiliser. Ammonium sulfate solution can be prepared in the laboratory using dilute ammonia solution and dilute sulfuric acid.

Describe a titration experiment to prepare ammonium sulfate solution using  $25.0 \text{ cm}^3$  of dilute ammonia solution, then to produce pure, dry crystals from this solution. (6 marks)

### Student answer

Clamp a burette vertically and fill it with dilute sulfuric acid. Run out enough acid to remove any air from the tip. Record the burette reading from the bottom of the meniscus. Use a pipette with a pipette filler to transfer  $25.0 \text{ cm}^3$  of the ammonia solution to a conical flask, and add a few drops of phenolphthalein indicator [1]. Place the conical flask on a white tile underneath the burette.

Add acid from the burette, swirling the flask gently. Add the acid drop by drop near the end-point until the indicator just changes from pink to colourless [2]. Record the burette reading and calculate the titre (end reading minus start reading). Add the titre volume of acid to  $25.0 \text{ cm}^3$  of ammonia solution, but this time without the indicator [3].

Pour the mixture into an evaporating basin and heat over a boiling water bath to produce a concentrated solution. Allow the solution to cool and crystals to form. Remove the crystals and dry them in a warm oven [4].

[1] This is a good answer as it names a specific indicator (though methyl orange indicator would be a much better choice because it is more suited to titrations with strong acids and weak alkalis like ammonia solution).

[2] This is an important step to obtain an accurate titre and so it is important to include it in the answer. The answer could also describe washing the insides of the flask with distilled water near to the end-point.

[3] It would be better to carry out the experiment more than once to obtain concordant results (identical or very similar titres).

[4] The answer includes how to produce pure, dry crystals from the ammonium sulfate solution – a part of the answer that would be easy to forget to write.

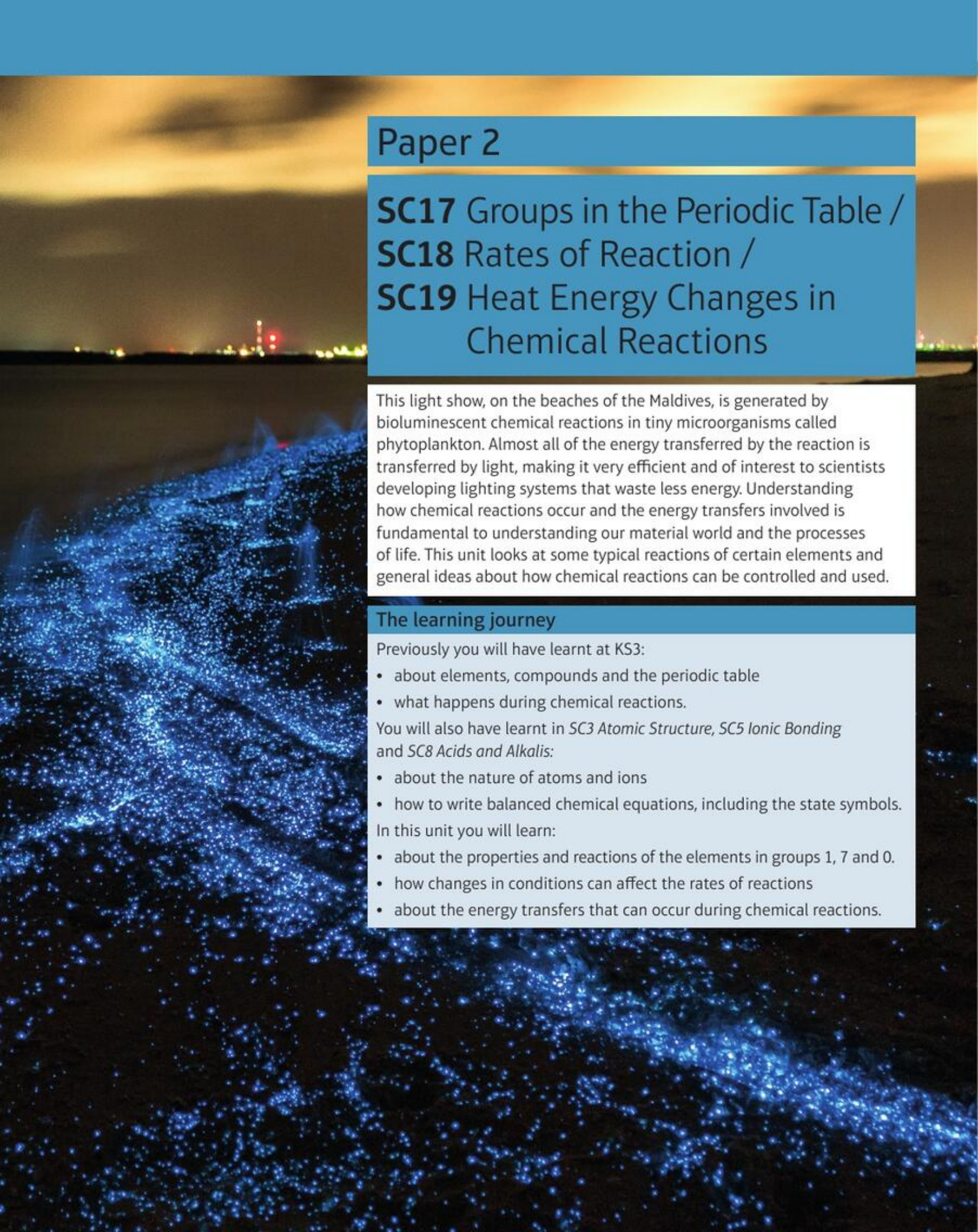
### Verdict

This is a strong answer. It shows good knowledge and understanding of how to carry out a titration and how to use crystallisation to produce pure, dry crystals. The answer describes the steps needed in a logical order. It gives practical details including the expected colour change at the end-point of the titration, and apparatus needed for safe working such as a pipette filler.

#### Exam tip

The specification contains a list of practicals that you should have covered in lessons. Make sure you have revised the practicals and can name the pieces of apparatus involved.





## Paper 2

### **SC17** Groups in the Periodic Table / **SC18** Rates of Reaction / **SC19** Heat Energy Changes in Chemical Reactions

This light show, on the beaches of the Maldives, is generated by bioluminescent chemical reactions in tiny microorganisms called phytoplankton. Almost all of the energy transferred by the reaction is transferred by light, making it very efficient and of interest to scientists developing lighting systems that waste less energy. Understanding how chemical reactions occur and the energy transfers involved is fundamental to understanding our material world and the processes of life. This unit looks at some typical reactions of certain elements and general ideas about how chemical reactions can be controlled and used.

#### The learning journey

Previously you will have learnt at KS3:

- about elements, compounds and the periodic table
- what happens during chemical reactions.

You will also have learnt in *SC3 Atomic Structure*, *SC5 Ionic Bonding* and *SC8 Acids and Alkalis*:

- about the nature of atoms and ions
- how to write balanced chemical equations, including the state symbols.

In this unit you will learn:

- about the properties and reactions of the elements in groups 1, 7 and 0.
- how changes in conditions can affect the rates of reactions
- about the energy transfers that can occur during chemical reactions.



# SC17a Group 1

Specification reference: C6.1; C6.2; C6.3; C6.4; C6.5

## Progression questions

- What are the main properties of alkali metals?
- How do alkali metals react with water?
- Why do alkali metals have different reactivities?

alkali metals		main group numbers										noble gases					
1	2											3	4	5	6	7	0
Li	Be											B	C	N	O	F	He
Na	Mg											Al	Si	P	S	Cl	Ar
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe
Cs	Ba	La	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn
Fr	Ra	Ac	Rf	Db	Sg	Bh	Hs	Mt	Ds	Rg							

The **periodic table** is arranged so that elements in the same vertical column or **group** have similar chemical and physical properties, and show trends in those properties.

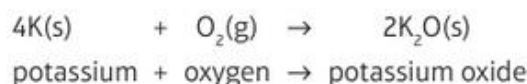
The **alkali metals**, in group 1, have similar physical properties to other metals – they are all malleable and conduct electricity. However, they also have properties that are specific to this group. All alkali metals have relatively low melting points, are soft and easily cut. Alkali metals are also very reactive and readily form compounds with non-metals.

**A** Groups 1, 7 and 0 in the periodic table have special names.



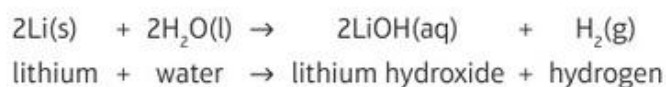
- 1** Why are potassium and sodium placed in the same group of the periodic table?

All alkali metals are easily oxidised and burn brightly in air. For example, the reaction of potassium with oxygen can be shown as:



- 2** Describe two physical properties of alkali metals that make them different from other metals.

The reactions of three alkali metals with water are described in table B. In each reaction, a metal hydroxide (an alkali) and hydrogen gas are the products. The **reactivity** of the alkali metals increases down the group.



- 3** Write word equations for the reactions of sodium with:

- oxygen
- water.

reactivity ↓	lithium + water	bubbles fiercely on the surface
	sodium + water	melts into a ball and fizzes about the surface
	potassium + water	bursts into flames and flies about the surface



**B** reactions of alkali metals with water



- 4** Caesium (Cs) is below potassium in the periodic table. Suggest how caesium might react with water.

## Did you know?

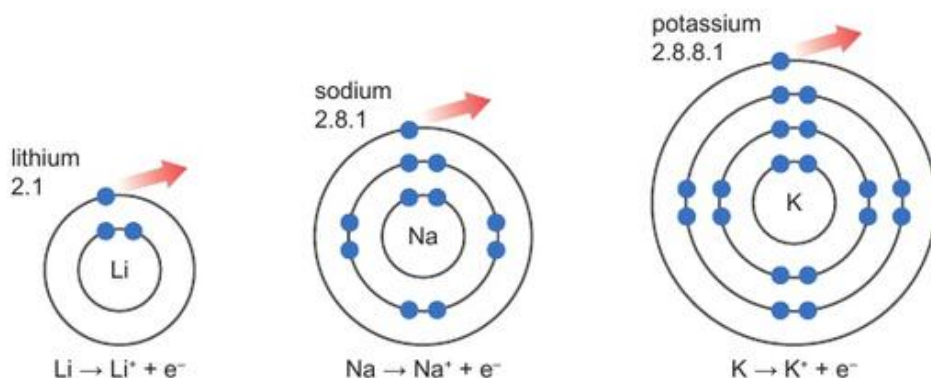
Francium, discovered in 1939, was the last element to be discovered in nature. It is radioactive and extremely rare. Scientists think that only 20–30g of francium exists on Earth at any one time.



To explain the trend in reactivity we need to look at the electronic configurations of alkali metal atoms. These atoms have one electron in their outer shells. When they react with non-metal atoms, the outer electrons are transferred from the metal to the non-metal. Diagram D shows an example. Each sodium atom loses its outer electron to form a positive (1+) ion, and each oxygen atom gains two electrons to form a negative (2-) ion.

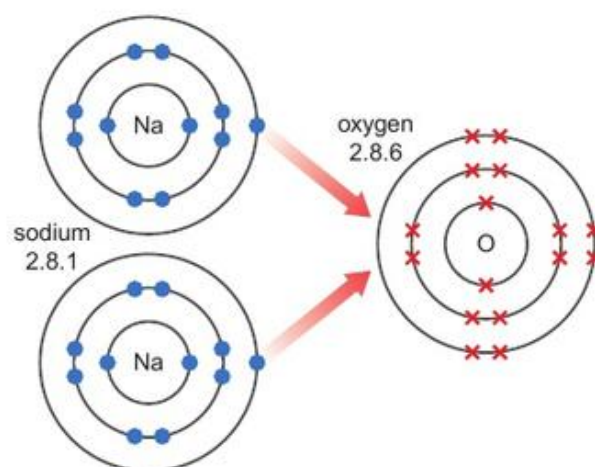
- 5** Draw diagrams, like the ones in diagram C, to show the electronic configurations of the sodium and oxide ions.

As we go down group 1, the atoms get larger because an extra electron shell is added in each period. The force of attraction between the positive nucleus and the negative outer electron decreases as they become further apart. This explains the trend in reactivity of alkali metals. Sodium is more reactive than lithium because it is easier to remove the outer electron from a sodium atom.



**D** As the distance between the outer electron and the nucleus increases, the alkali metals get more reactive.

- 8** By referring to atomic structures, explain why potassium is more reactive than sodium.
- 9** **H** Write an ionic equation for the reaction of potassium with water (hydroxide ion = OH<sup>-</sup>).



**C** dot and cross diagrams to show sodium reacting with oxygen to form sodium oxide (Na<sub>2</sub>O, which contains Na<sup>+</sup> and O<sup>2-</sup> ions)

- 6** Why do alkali metals form an ion with a 1+ charge?
- 7 a** Why are alkali metals stored under oil?
- b** How will the reactivity of rubidium (Rb) compare to potassium and caesium?

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Name three alkali metals and describe their main physical and chemical properties.

### Extend

- E1 a** Explain the difference in reactivity between rubidium and caesium.
- b** **H** Write ionic equations for the reactions of rubidium and caesium with water.

### Exam-style question

- a** Write a balanced equation for the reaction of potassium with water. (2 marks)
- b** Explain why the reaction of caesium with water is not demonstrated in school laboratories. (1 mark)

# SC17b Group 7

Specification reference: C6.6; C6.7; C6.8; C6.9; C6.10

## Progression questions

- How do the physical properties of the halogens change, going down group 7?
- How can we test for chlorine gas?
- How do halogens react with metals and hydrogen?



A the three most common halogens

Elements in group 7 of the periodic table are called the **halogens**. They all share similar properties and show a pattern in the way their properties change through the group.

All halogens exist as **diatomic** molecules, with two atoms held together by a single covalent bond. They are all non-metallic elements, which are poor conductors of heat and electricity. Care has to be taken when handling halogens, as they are all toxic and corrosive.



**1** What pattern is there in the depth of colour of the halogens?

The physical properties of the three most common halogens are described in table B. As you go down the group the melting points, boiling points and densities increase.

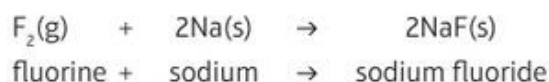
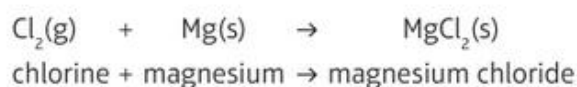
## Did you know?

The name halogen comes from the Greek words *hals*, meaning 'salt', and *gen*, meaning 'to make'. So halogens make salts – not just table salt (sodium chloride) but a range of metal halides that have a variety of uses.

Halogen	Melting point (°C)	Boiling point (°C)	Density (g/cm <sup>3</sup> )	Appearance
chlorine	-101	-34	0.0032	green gas
bromine	-7	59	3.12	brown liquid
iodine	114	184	4.95	purple/black solid

B halogen properties

Most halogens react with metals and non-metals in a similar way. Halogens react with metals forming ionic compounds called **salts**, which contain **halide** ions (X<sup>-</sup>). For example:



Halide salts have many uses. For example, common sodium halide salts include sodium chloride (table salt), sodium fluoride (found in many toothpastes), sodium bromide (used in a disinfectant for swimming pools) and sodium iodide (added to table salt to prevent iodine deficiency).



**2** Describe two ways that halogens are similar.

**3** Look at photo A and table B.



**a** What do you think fluorine, at the top of group 7, will look like?



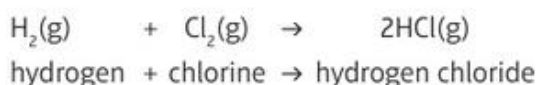
**b** Estimate the melting point, boiling point and density of fluorine.



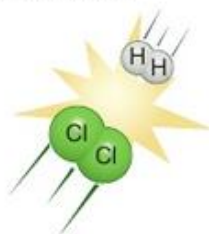
- 4 What ions are formed when sodium reacts with fluorine?
- 5 a Name the product formed when calcium metal burns in fluorine.  
b Write a balanced equation for the reaction (calcium ion =  $\text{Ca}^{2+}$ ).

All halogens can be used as **disinfectants** and **bleaches**, as they can kill microorganisms and remove the colour from materials. Chlorine is commonly used in swimming pools and many types of bleach. The test for chlorine, shown in photo C, depends on this bleaching action.

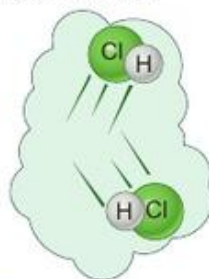
Halogens react with hydrogen to form hydrogen halides, which dissolve in water to form acidic solutions. For example, hydrogen and chlorine explode to form hydrogen chloride, which dissolves in water to make hydrochloric acid.



Hydrogen and chlorine molecules collide and the covalent bonds holding the atoms together break.

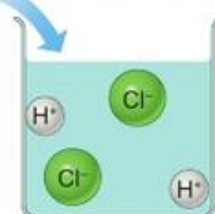


Covalent bonds form between hydrogen and chlorine atoms, making a new compound, hydrogen chloride.



When hydrogen chloride dissolves in water the molecules break up into two ions,  $\text{H}^+$  and  $\text{Cl}^-$ .

The formation of  $\text{H}^+$  ions makes the solution acidic.



dissolves in water to form hydrochloric acid

D making hydrochloric acid

- 7 In what state would each of the halogens in table B be at:
- a  $-50^\circ\text{C}$       b  $150^\circ\text{C}$ ?
- 8 Write word equations for the reactions between:
- a lithium and chlorine      b hydrogen and fluorine.
- 9 Explain how hydrogen fluoride can form an acidic aqueous solution and name the acid.
- 10 Write balanced equations for the reactions between:
- a sodium and iodine      b hydrogen and bromine.

### Exam-style question

- a Predict the state and appearance of astatine (At). (1 mark)  
b Explain your answer. (1 mark)



C If damp blue litmus paper is placed in chlorine gas, it first turns red then bleaches white.

6 Describe for chlorine:

- a two uses  
b a safety precaution for its use  
c a test for the gas.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

S1 Describe how the halogens chlorine, bromine and iodine are similar, and how they show trends in properties down their group.

### Extend

E1 Describe, using balanced equations where appropriate, how bromine can be converted into:

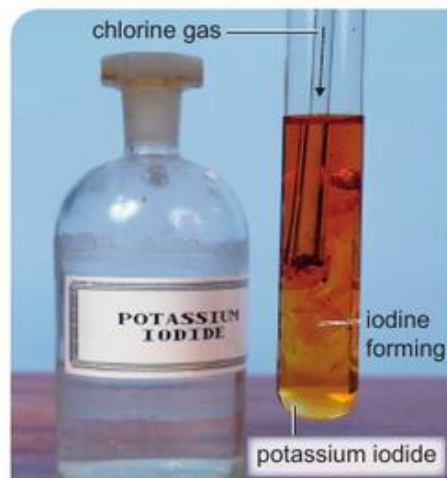
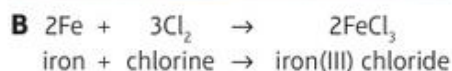
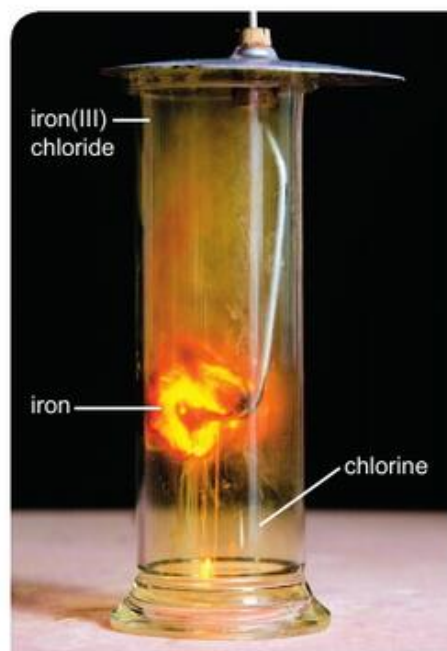
- a sodium bromide  
b hydrobromic acid.

# SC17c Halogen reactivity

Specification reference: **H** C0.4; C6.11; **H** C6.12; C6.13

## Progression questions

- How can displacement reactions be used to work out the reactivity of halogens?
- How can we explain the reactivity of halogens?
- **H** What happens to halogen atoms and halide ions during displacement?



**C** Chlorine displaces iodine from potassium iodide solution.

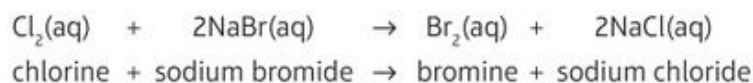
Table A shows the order of reactivity of different halogens when heated with iron wool. In general the halogens become less reactive as you go down the group.

Halogen	Effect on iron wool
fluorine	bursts into flames
chlorine	glows brightly
bromine	glows dull red
iodine	changes colour

**A** halogens and heated iron wool

- 1** Astatine, symbol At, is found below iodine in the halogen group.
  - a** Write a word equation for the reaction of astatine with iron wool.
  - b** How will this reaction compare with chlorine? Explain your answer.
- 2** Write a balanced equation for the formation of magnesium iodide ( $\text{MgI}_2$ ) from its elements.

The order of reactivity of halogens can also be worked out using **displacement reactions**. In a displacement reaction, a more reactive element takes the place of a less reactive element in an ionic compound. So, a more reactive halogen displaces a less reactive halogen from a halide compound. For example, chlorine displaces bromine from sodium bromide in solution, but bromine cannot displace chlorine from sodium chloride.

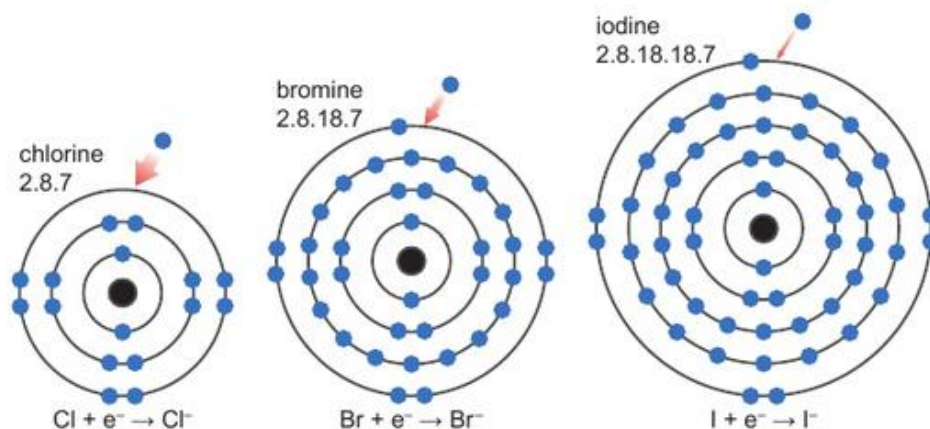


- 3** Which of the following pairs of substances do not react? Explain your choice.
  - a**  $\text{Br}_2 + \text{LiCl}$
  - b**  $\text{Cl}_2 + \text{NaI}$
- 4** Write a balanced equation, with state symbols, for the reaction that occurs in photo C.
- 5** What would be observed if bromine gas were bubbled through potassium iodide solution?



To explain the trend in reactivity we need to look at the electronic configuration of the halogen atoms.

Group 7 atoms gain one electron when they react. Down the group, the distance between the outermost shell containing electrons and the nucleus increases. This means that the force of attraction between the positive nucleus and an incoming negative electron decreases, and so the ions do not form so easily and the reactivity decreases.



**D** Going down group 7, the outermost electron shell gets further from the nucleus and the ions are less readily formed.



**6** Explain why fluorine is the most reactive halogen and astatine is the least reactive halogen.

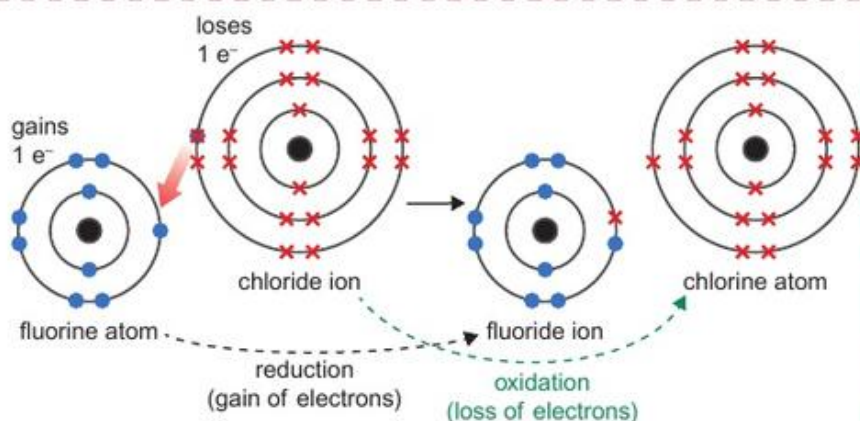
## H Redox

When a metal reacts with oxygen it loses electrons and so we can define **oxidation** as a 'loss of electrons'.

**Reduction** is the opposite and is a 'gain of electrons'. Use the mnemonic 'OILRIG' to remember that 'Oxidation Is Loss, Reduction Is Gain'. These two processes occur at the same time in displacement reactions, which makes them examples of reduction-oxidation or **redox** reactions.

Diagram E shows the reaction between fluorine atoms and chloride ions.

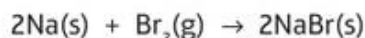
As the fluorine atoms are **reduced**, by gaining electrons, the chloride ions are **oxidised**, by losing electrons.



**E** Fluorine gains electrons more readily than chlorine. It displaces chlorine from chloride ions:  $\text{F}_2(\text{g}) + 2\text{Cl}^-(\text{aq}) \rightarrow 2\text{F}^-(\text{aq}) + \text{Cl}_2(\text{g})$

**7 a** What happens in all redox reactions?

**b** Identify the substances that are oxidised and reduced in the reaction below. Explain your choice.



**8** Write an ionic equation with state symbols for the reaction that occurs in question 5.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** Design a summary table or diagram to describe and explain the trend in reactivity of the halogens.

## Extend

**E1 H** Use a displacement reaction to explain what is meant by 'redox reactions'.

## Exam-style question

Compare the trends in melting point and reactivity of group 1 and group 7 elements. (3 marks)

The noble gases have a variety of uses because of their unique properties.

Krypton is used in photography lighting. It produces a brilliant white light when electricity is passed through it.

Argon is denser than air. It is added to the space above the wine in wine barrels to stop oxygen in the air reacting with the wine.

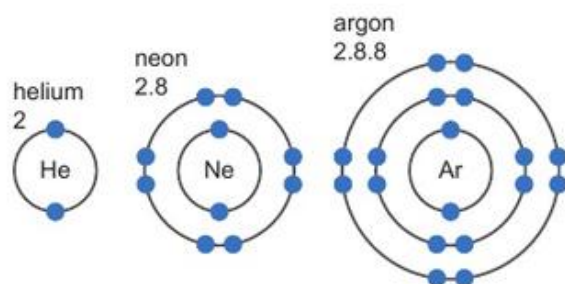
Helium has a very low density and is non-flammable, so it is used in weather balloons and airships.

Neon produces a distinctive red-orange light when electricity is passed through it. This property makes it useful for making long-lasting illuminated signs.



**D** helium-filled balloons

To explain why noble gases are unreactive, we need to look at their electronic configurations. Diagram F shows that all noble gases have a complete outer shell of electrons. It has been shown that, when atoms form bonds, they can become more stable. They do this by gaining, losing or sharing electrons to get an electronic configuration like a noble gas. So noble gases are unreactive because their atoms already have a stable electronic configuration with a complete outer shell.



**F** Noble gases do not react as they already have a complete outer shell of electrons.

- 7** Explain why argon is very unreactive while potassium, with one more electron, is very reactive.

### Exam-style question

Neon is a colourless inert gas.

- a** Describe what the term inert tells you about neon. (1 mark)  
**b** Explain why neon exists as single atoms. (2 marks)

### Did you know?

Helium atoms are continually being produced by radioactive decay in certain rocks. However, the atoms are so light that most of them escape into space as the Earth's gravity cannot hold them.



**E** a neon-filled advertising sign

- 6 a** Explain how the properties of helium gas make it useful for weather balloons.  
**b** Explain why argon can be used in fire extinguishers.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Explain how the electronic configuration of noble gases affects their properties.

### Extend

- E1** The noble gas radon is radioactive. Use the information on these two pages to predict other properties of radon.



# SC18a Rates of reaction

Specification reference: C7.2; C7.5

## Progression questions

- What changes can occur as a reaction proceeds?
- How can we investigate rates of reaction?
- How are graphs used to show rates of reaction?



**A** The combustion of the wood is a faster reaction than the baking of the pizzas.

New substances are always formed in chemical reactions. The **rate** of a reaction is the speed at which **reactants** are turned into **products**. Rates vary greatly, from very slow (e.g. the rusting of iron) to almost instantaneous (e.g. explosions, precipitation reactions). However, we can control rates of reaction by altering **variables**, such as the concentration of solutions and the size of pieces of solid reactant.

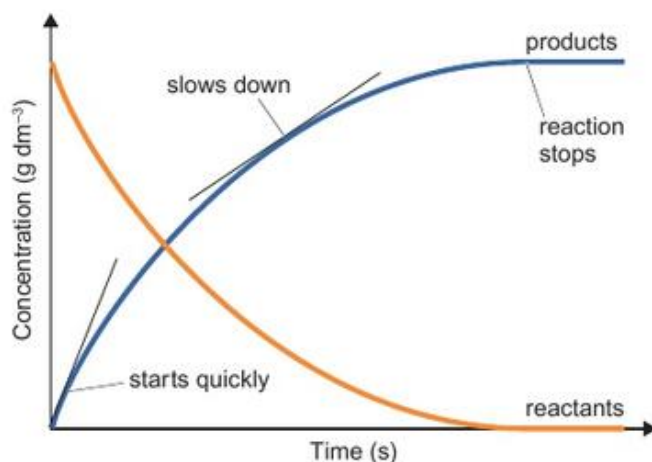


- 1** Describe one example of a very fast and a very slow chemical reaction.

To investigate reaction rates, we need to be able to measure how the amount of reactants or products changes with time. Graph B shows how the concentrations change during a reaction. The gradient (slope) of the graph indicates the rate: the steeper the graph, the faster the reaction. Reaction rates are usually fastest at the start because that is when the concentration of the reactants is greatest.

## Did you know?

Japanese scientists have developed a camera to observe extremely fast reactions. The camera records 4.4 trillion frames per second. A good smartphone would need over 1000 years to record as many frames as this camera records in the blink of an eye.



**B** Reactions are usually fastest at the start.



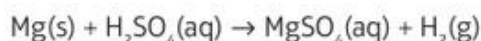
- 2** What happens to the concentration of reactants and products during a reaction?

We can also monitor rates by measuring changes in the mass or volume of reactants or products.

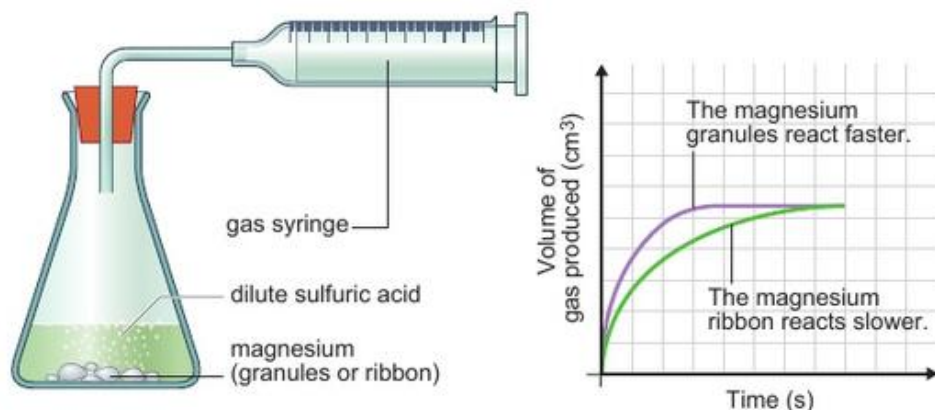


- 3** Why do most reactions slow down in time?

In the reaction between magnesium and sulfuric acid, shown in diagram C, we can follow the reaction by measuring the volume of hydrogen gas produced.



The results show the difference between using magnesium ribbon and magnesium granules. The graph for the granules is steeper and levels off more quickly. This shows that the granules, which are smaller pieces of solid than the ribbon, react more quickly.



**C** In this experiment only the size of the pieces of magnesium is changed. All other variables are kept constant.

In the reaction between marble (calcium carbonate) and hydrochloric acid, shown in diagram D, we can follow the rate by measuring the change in mass. The mass decreases as carbon dioxide gas escapes from the flask.



**6** Calcium carbonate is left at the end of the reaction shown in diagram D.



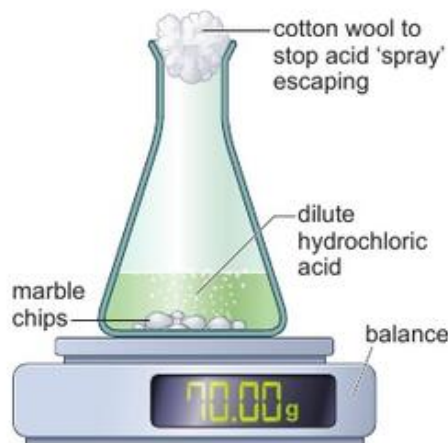
**a** Which reactant has been used up?



**b** How do you know when the reaction is complete?



**c** Sketch a graph to show how the mass of the flask will change as the reaction proceeds.



**D** As the reaction proceeds, the mass of the flask and contents will decrease.



**7** Why would it be difficult to measure the reaction rate for the rusting of iron?



**4 a** What piece of apparatus can be used to measure gas volumes accurately?



**b** Explain why measuring gas volume can be used to monitor the rates of some reactions.



**5** How does the graph in diagram C show that magnesium granules react faster than ribbon?

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Look at the reaction in diagram D.

- What is being measured in this experiment?
- What happens to the concentration of the acid as the reaction proceeds?
- What other change could be measured to follow this reaction?

### Extend

**E1** Describe, including diagrams, how you could use a gas syringe to investigate the correlation between temperature and the rate of the reaction between magnesium and hydrochloric acid.

### Exam-style question

When copper carbonate reacts with sulfuric acid, carbon dioxide gas is formed. Explain how an electronic balance could be used to investigate the rate of this reaction. (2 marks)

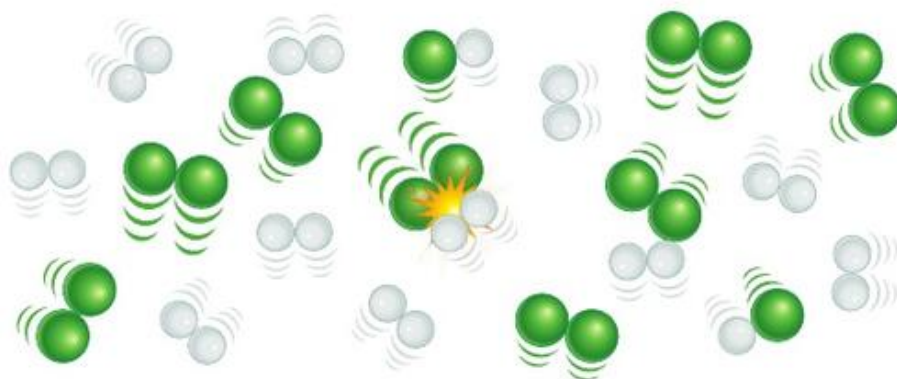


# SC18b Factors affecting reaction rates

Specification reference: C7.3; C7.4

## Progression questions

- What has to happen for two particles to react?
- How does the speed of particles affect the rate of reaction?
- Why do changes in temperature, concentration, surface area and pressure affect rates of reaction?



**A** The faster-moving hydrogen and chlorine molecules have more energy, and are more likely to react.

For chemical reactions to occur the reactant particles must collide or 'bump' together with enough energy to react. The minimum amount of energy needed for a reaction to occur is called its **activation energy**. During successful collisions, this energy helps to break bonds, so that the atoms can be rearranged to make new substances (the products of the reaction).



- 1** What two things must happen before hydrogen and chlorine can react?

When methane and air are mixed the molecules collide, but no reaction occurs until a spark or flame provides the activation energy needed. The reaction then keeps going because the reaction releases energy, which then provides the activation energy needed. Reactions which release energy are described as **exothermic** changes.



**B** The natural methane leaking out of these rocks has been burning for decades, having been set alight by lightning.



- 2** Why does a Bunsen burner not light as soon as you turn on the gas?



- 3** What is the activation energy of a reaction?



- 4** State two changes that would make the reaction in diagram A slower.

A few chemical reactions need to take in energy to occur. These reactions, called **endothermic** changes, will not keep going unless energy is continually supplied. For example, the electrolysis of copper chloride needs a continuous supply of electrical energy.

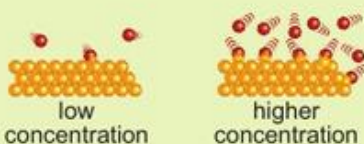
In general, reaction rates are increased when the energy of the collisions is increased, and when the frequency is increased (the number of collisions in a certain amount of time). More collisions occur if the particles are closer together or moving faster. The particles in any substance have a range of energies, but only those with enough energy can react. More collisions will be successful if more of the particles have the activation energy required.



**Concentration and reaction rate**

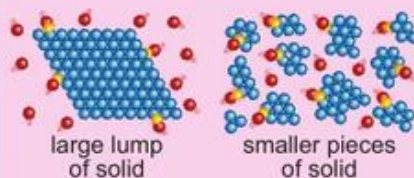
*Change:* Increasing the concentration of solutions increases the rate of reaction.

*Explanation:* There are more reacting particles in the same volume so collisions occur more often.

**C****Surface area and reaction rate**

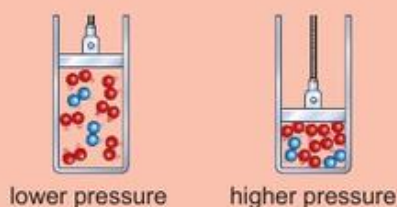
*Change:* Increasing the surface area to volume ratio, by decreasing the size of solid pieces while keeping the total volume of solid the same, increases the rate of the reaction.

*Explanation:* There is more surface for collisions to occur on, so collisions occur more often.

**D****Pressure of gases and reaction rate**

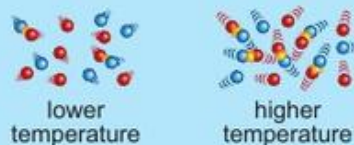
*Change:* Increasing the pressure of gases increases the rate of reaction.

*Explanation:* The reactant particles are squeezed closer together so collisions occur more often.

**E****Temperature and reaction rate**

*Change:* Increasing the temperature increases the rate of reaction.

*Explanation:* The reactant particles speed up and have more energy. They therefore collide more often and more particles have enough energy to react when they collide.

**F**

- 7** Suggest a reason why reactions between gases are often slower than reactions between liquids at the same temperature.

- 5 a** Which of the following burns most quickly in air: wood chips, wood dust or wood shavings?

- b** Explain your answer to part a.

- c** Explain how using pure oxygen instead of air would affect the rate of burning.

- 6** The reaction between hydrogen ( $H_2$ ) and chlorine ( $Cl_2$ ) gases produces hydrogen chloride ( $HCl$ ) gas. For this reaction:

- a** write a balanced chemical equation, with state symbols

- b** explain how and why decreasing the gas pressures affects the rate

- c** explain how and why increasing the temperature affects the rate.

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

- S1** Use ideas about collisions and energy to explain how the rate of a reaction can be increased by changes in concentration, temperature and the size of pieces of a solid reactant.

**Extend**

- E1** Explain four ways in which the rate of reaction between iron lumps and oxygen from the air can be increased.

**Exam-style question**

Explain why acid reacts faster with powdered chalk than with lumps of chalk. (3 marks)



# SC18b Core practical – Investigating reaction rates

Specification reference: C7.1

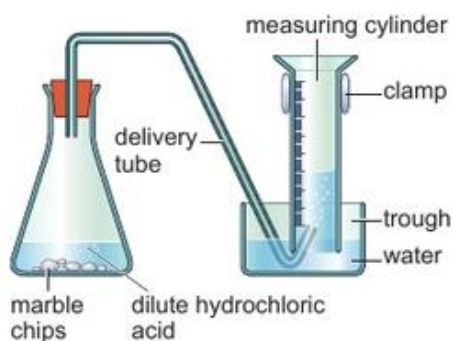
## Aim

Investigate the effects of changing the conditions of a reaction on the rates of chemical reactions.

The progress of a chemical reaction can be measured by how the amounts of reactant or product change with time, or by the time taken for the reaction to reach a certain point.

## Your task 1 – measuring volumes of gases

You are going to investigate the reaction between hydrochloric acid and marble chips (calcium carbonate) and how the surface area of the marble chips affects the rate. You will monitor the progress of the reaction by measuring the volume of carbon dioxide produced.



**A** investigating volumes of gas produced

## Method 1

Wear eye protection.

- A** Set up the apparatus as shown in diagram A.
- B** Measure 40 cm<sup>3</sup> of dilute hydrochloric acid into a conical flask.
- C** Add 5 g of small marble chips to the flask.
- D** Immediately stopper the flask and start the stop clock.
- E** Note the total volume of gas produced after every 30 seconds until the reaction has finished.
- F** Repeat the experiment using 5 g of larger marble chips.

## Your task 2 – observing a colour change

You are going to investigate the effect of temperature on the rate of reaction between sodium thiosulfate and hydrochloric acid. You will monitor the progress of the reaction by observing a colour change (as shown in photo B).



**B** We can follow the rate of the reaction between sodium thiosulfate and hydrochloric acid by measuring the time taken for a 'cross' drawn beneath the reaction beaker to disappear.

## Method 2

Wear eye protection.

- G** Place 50 cm<sup>3</sup> of sodium thiosulfate solution into a 300 cm<sup>3</sup> conical flask.
- H** Measure out 5 cm<sup>3</sup> of dilute hydrochloric acid in a test tube.
- I** Clamp the conical flask in place in a water bath at a certain temperature. Place the test tube in a rack in the same water bath. Record this temperature.
- J** After 5 minutes, remove the flask and place it on a piece of white paper marked with a cross.
- K** Add the acid to the thiosulfate and start the stop clock.

- L** Looking down from above, stop the clock when the cross disappears.  
**M** Note this time and take the final temperature of the mixture.  
**N** Repeat at three or four other temperatures, between 20 and 50 °C.

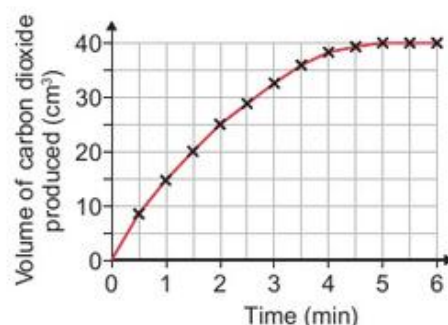
### Exam-style questions

Look at Method 1.

- 1** Write a word equation and a balanced symbol equation for this reaction. (4 marks)
- 2 a** State the dependent and independent variables in this investigation. (2 marks)
- b** State two control variables. (2 marks)
- 3 a** State why you must immediately stopper the flask in step D. (1 mark)
- b** State the apparatus you would use to measure the mass of the marble chips. (1 mark)
- 4** Look at graph C.
  - a** State when the reaction is complete. Explain your answer. (2 marks)
  - b** Sketch graph C and add a curve that would be produced by smaller marble chips. (2 marks)
  - c** Use graph C to calculate the average reaction rate in  $\text{cm}^3/\text{min}$  between 45 and 105 seconds. Show your working. (2 marks)
  - d** Describe how you would use a tangent line to estimate the reaction rate in  $\text{cm}^3/\text{min}$  at 100 seconds. (3 marks)
- 5** Describe how you would modify Method 1 to investigate the effect of temperature on the rate of this reaction.
- 6** State one further variable that would effect the rate of this reaction. (1 mark)

Look at Method 2.

- 7** State why the investigation would be improved by measuring the initial temperature of the reaction mixture (just after the acid is added) and then measuring the final temperature (when the reaction is finished). (2 marks)
- 8** Some results from the experiment in method 2 are shown in table D.
  - a** State why the cross disappears. (1 mark)
  - b** Sketch a graph of the results (no graph paper required) with temperature on the horizontal axis. (3 marks)
  - c** Explain what these results tell us about the effect of temperature on the rate of this reaction. (2 marks)
  - d** Describe one way of improving the results obtained from this investigation. (1 mark)



**C** volume of gas collected in the reaction between hydrochloric acid and marble chips

Average temperature (°C)	Time for cross to disappear (s)
20	165
30	81
40	42
50	21

**D**

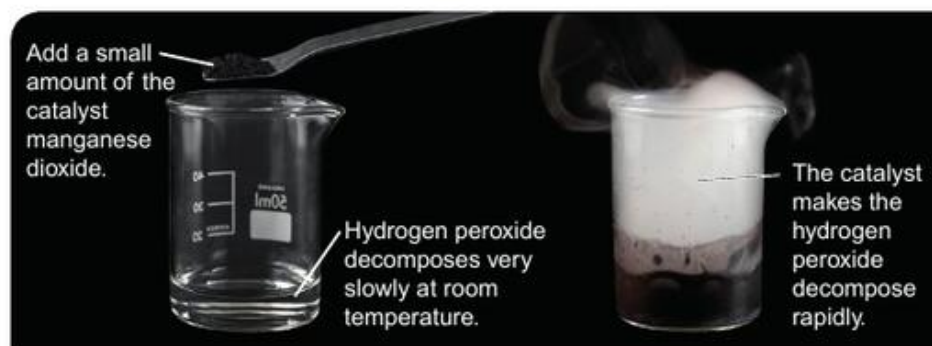


# SC18c Catalysts and activation energy

Specification reference: C7.6; C7.7; C7.8

## Progression questions

- What is a catalyst?
- How do catalysts work?
- What are enzymes used for?



**Catalysts** are substances that speed up chemical reactions without being permanently changed themselves and without altering the products of the reaction. Photo A shows the effect of a catalyst (manganese dioxide) on the decomposition of hydrogen peroxide.

**A** The catalyst is not used up. So, if 1 g of manganese dioxide is added at the start, then 1 g will be recovered at the end.

Catalysts are often used in industry. For example, platinum is used as a catalyst in the manufacture of nitric acid from ammonia. These catalysts make industrial processes more profitable, by making products more quickly, and by allowing reactions to occur at lower temperatures, which saves costs. Catalysts also do not usually need to be replaced because they are not used up.

We can think about catalysts by using an analogy. In diagram B, the starting point and the finishing point are the same for all cars. However, those following the scenic route need more energy at the start to get over the hill. So, in terms of energy, most cars will use the road through the tunnel, which is the easier route.



**B** Not all cars have enough energy to go over the hill.

Like the tunnel, a catalyst provides an alternative reaction route, which requires less activation energy. **Reaction profiles** like graph C can be used to show the energy changes for a catalysed and an uncatalysed exothermic reaction. The catalyst does not alter the overall energy change. However, since less energy is needed to start the reaction, more reactant molecules have enough energy and so more collisions are successful. This means that the reaction is faster.

## Did you know?

Many washing powders contain biological catalysts called enzymes to help break down different types of stains: proteases break down proteins like blood and egg, amylases break down starches and lipases break down fats and grease.

- 1** Explain how catalysts reduce the costs in industrial processes.
- 2** Platinum is a very expensive catalyst. Why might this not matter?
- 3** How could you show that the manganese dioxide in photo A was not used up?

- 4 Explain whether a reaction with a high or low activation energy would be faster.
- 5 In terms of energy changes, what is the same for catalysed and uncatalysed reactions?

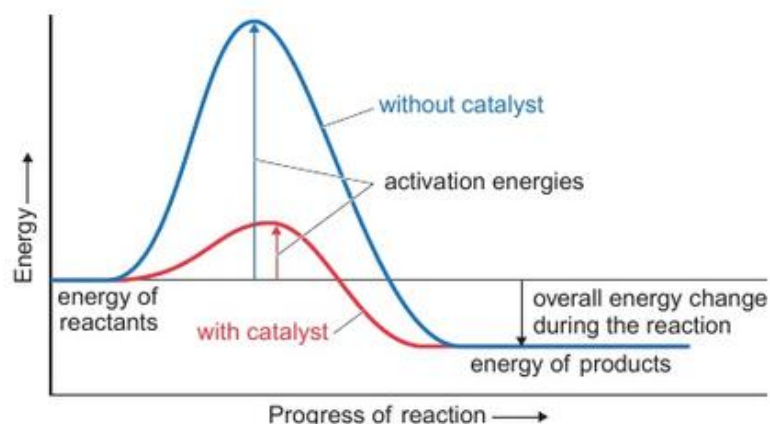
In catalytic converters in car exhausts (shown in diagram D) the catalysts are platinum and palladium. Their presence lowers the activation energy needed to convert harmful gases into harmless gases. The metals used are expensive but don't need to be replaced as they are not used up.

- 6 Explain why the iron catalyst used in the manufacture of ammonia:
- is in the form of small lumps
  - does not need to be replaced often.

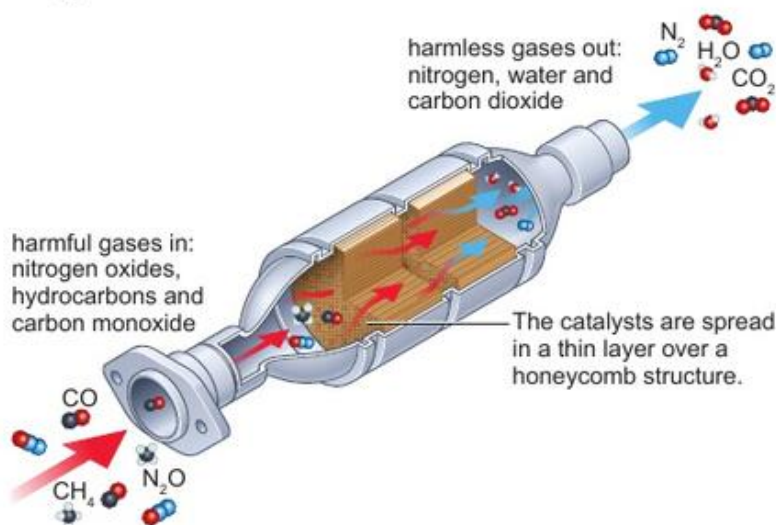
**Enzymes** are large complex **protein** molecules that act as catalysts in biological reactions. Each enzyme molecule has a part with a specific shape called the **active site**. The reactant molecules (**substrates**) fit the shape of the active site, a bit like a 'lock and key'. So each enzyme only fits one substrate and only catalyses one specific reaction. Enzymes are sensitive to changes in temperature and pH as these can **denature** their molecules (make them change shape) so they will not work.

Enzymes are essential in living things and many industries. For example, alcoholic drinks are produced using an enzyme in yeast. This enzyme catalyses the reaction in which glucose is converted into ethanol (alcohol) and carbon dioxide.

- 7 a Give an example of an enzyme and the reaction it catalyses.
- b Why will the enzyme not catalyse other reactions?
- 8 Explain why heating a reaction involving a catalyst will usually make it faster but heating an enzyme-catalysed reaction can make it slower.



C This reaction profile shows that a catalyst lowers the activation energy.



D The honeycomb structure gives the catalyst a large surface area.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Explain, in terms of activation energy, how catalysts and enzymes work and why they are useful in industrial chemical reactions.

### Extend

**E1** Compare and contrast chemical catalysts and biological catalysts.

### Exam-style question

Explain how the following speed up chemical reactions by increasing the frequency of successful collisions.

- using a catalyst (2 marks)
- increasing the temperature (2 marks)



# SC19a Exothermic and endothermic reactions

Specification reference: C7.9; C7.10; C7.11

## Progression questions

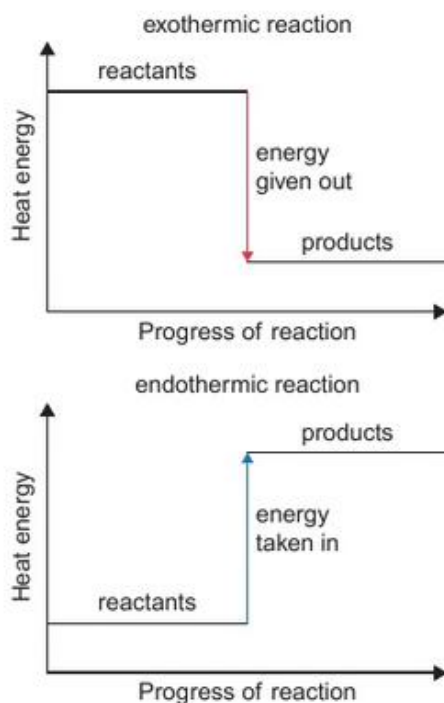
- What are exothermic and endothermic reactions?
- What are some examples of exothermic and endothermic reactions?
- How can heat changes in solution be investigated?



**A** This self-heating can uses an exothermic reaction between sodium hydroxide pellets and water.



- 1** Explain why combustion reactions are described as being exothermic.



**C** simple reaction profiles

Energy is transferred between the surroundings and the reactants during chemical reactions. Energy is most often transferred by heating. Chemical reactions can be described as:

- **exothermic** – energy is transferred from stores of energy in chemical bonds to the surroundings.
- **endothermic** – energy is transferred from the surroundings to stores of energy in chemical bonds.



**B** Combustion reactions are exothermic.

You can use a **reaction profile** to model the energy change during a chemical reaction. In these diagrams, energy stored in bonds is represented as a horizontal line. The greater the energy stored, the higher the line.



- 2** Explain how you can tell from its reaction profile whether a reaction is exothermic or endothermic.




- 3** Sherbet sweets contain citric acid and sodium hydrogen carbonate. These react together to produce sodium citrate, water and carbon dioxide. The reaction makes your mouth feel cold. Draw a reaction profile for this reaction, labelling each horizontal line with the names of the substances involved.

## Temperature changes during reactions

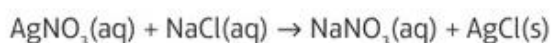
During a chemical reaction, energy is transferred between the reacting substances and their surroundings. This is usually by heating, particularly if a reaction takes place in solution. The stored thermal (heat) energy in the solution increases during an exothermic reaction, and it decreases during an endothermic reaction. This means you can determine whether a reaction in solution is exothermic or endothermic:

- the temperature increases in an exothermic reaction
- the temperature decreases in an endothermic reaction.

Diagram D shows a simple apparatus for these investigations.

-  **4** Explain why a lid and a *polystyrene* cup are used when investigating temperature changes in solution.


In **precipitation** reactions, an insoluble product forms from two solutions. For example:



Precipitation reactions can be exothermic or endothermic, depending on the substances involved. However, two types of reactions are always exothermic:

- **neutralisation**, the reaction between an acid and a base
- **displacement**, the reaction between a metal and a compound of a less reactive metal, or between a halogen and a compound of a less reactive halogen.

When a salt dissolves in water, the change is exothermic or endothermic, depending on the salt. The temperature increases as calcium chloride dissolves, but it decreases when ammonium chloride dissolves.

-  **5** Explain the temperature change you would expect when sodium hydroxide dissolves in water.

### Exam-style question

Sodium carbonate solution reacts with calcium chloride solution. A precipitate of calcium carbonate forms. A student carries out an experiment to measure the temperature change in this reaction. These are her results:

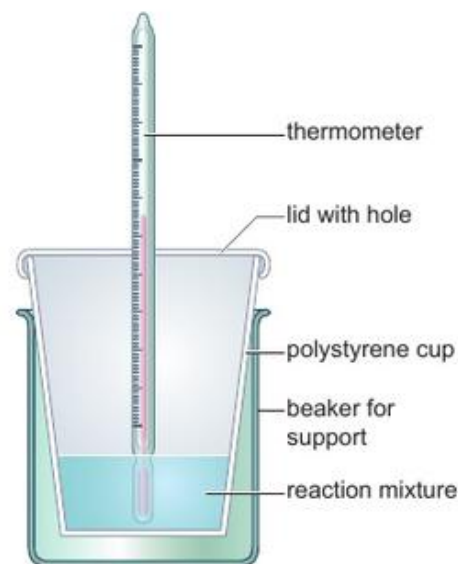
temperature of reactants = 21 °C

final temperature of mixture = 12 °C

Explain what this shows about the type of energy change that happens.  
(2 marks)

### Did you know?

Photosynthesis is an endothermic reaction.



**D** The air trapped between the beaker and the cup reduces energy transfers by heating, to and from the surroundings.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** How can you work out whether a reaction in solution is exothermic or endothermic?

### Extend

**E1** The temperature changes in displacement reactions between metals and solutions of metal compounds can be used to determine a reactivity series. Explain which variables should be controlled to obtain valid results.



# SC19b Energy changes in reactions

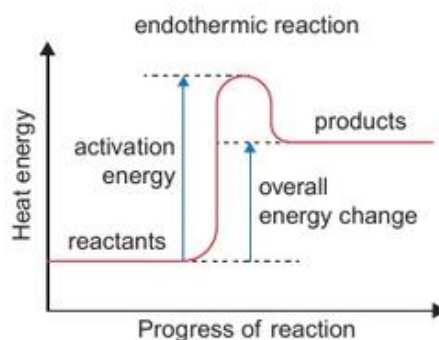
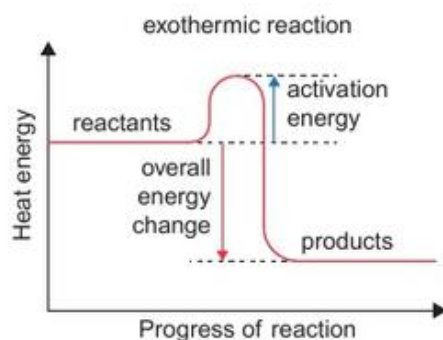
Specification reference: C7.12; C7.13; **H** C7.14; C7.15; C7.16

## Progression questions

- How can exothermic and endothermic reactions be explained in terms of bonds?
- How are exothermic and endothermic reactions modelled?
- **H** How are energy changes in reactions calculated?



**A** These bikes are fuelled with pure methanol.



**B** The activation energy is the difference in energy between the reactants and the top of the 'hump'.

Reactions happen when reactant particles collide with each other with sufficient energy. The **activation energy** is the minimum amount of energy needed by colliding particles for a reaction to happen. Precipitation and neutralisation reactions have low activation energies, and start as soon as the reactants are mixed. Combustion reactions have higher activation energies.



- 1** Explain why methanol needs a flame to start it burning.

You can model activation energy in a **reaction profile** by drawing a 'hump' between the reactants and products lines. Some reactions, such as combustion, have high activation energies and need energy from a spark or flame to start.



- 2** Describe how the activation energy of a reaction is modelled on a reaction profile.



- 3** Explain, in terms of bond breaking and bond making, why the combustion of methanol is an exothermic reaction.

## Breaking and making bonds

During a chemical reaction, bonds in the reactants break and new bonds are made to form the products:

- energy is transferred to the reactants to break their bonds, so breaking bonds is endothermic
- energy is transferred to the surroundings as bonds form, so making bonds is exothermic.

A reaction is exothermic if more energy is given out making bonds than is needed to break bonds. A reaction is endothermic if less energy is given out making bonds than is needed to break bonds.

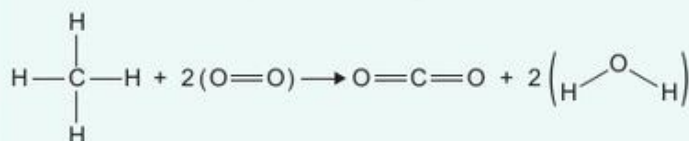
## H Bond energy calculations

A **mole** of something is  $6.02 \times 10^{23}$  of them (see SC9c Moles). The energy needed to break one mole of a particular **covalent bond** is its **bond energy**, measured in kilojoules per mole ( $\text{kJ/mol}$  or  $\text{kJ mol}^{-1}$ ). For example, the bond energy of a C–O bond is  $358 \text{ kJ mol}^{-1}$ . This means that 358 kJ must be taken in to break one mole of C–O bonds, and 358 kJ is given out when one mole of C–O bonds is made. Table C shows some bond energies.

You can use bond energies to calculate the energy change in a reaction, as shown in the worked example below.

### Worked example

Methane burns completely in oxygen to form carbon dioxide and water:



#### D

Calculate the energy change during this reaction.

#### Step 1 Calculate energy in (bonds broken)

$$\begin{array}{lll} 4 \times (\text{C}-\text{H}) & = 4 \times 413 & = 1652 \text{ kJ mol}^{-1} \\ 2 \times (\text{O}=\text{O}) & = 2 \times 498 & = 996 \text{ kJ mol}^{-1} \\ \text{Total in} & = 1652 + 996 & = 2648 \text{ kJ mol}^{-1} \end{array}$$

#### Step 2 Calculate energy out (bonds made)

$$\begin{array}{lll} 2 \times (\text{C}=\text{O}) & = 2 \times 805 & = 1610 \text{ kJ mol}^{-1} \\ 4 \times (\text{O}-\text{H}) & = 4 \times 464 & = 1856 \text{ kJ mol}^{-1} \\ \text{Total out} & = 1610 + 1856 & = 3466 \text{ kJ mol}^{-1} \end{array}$$

#### Step 3 Energy change = energy in – energy out

$$= 2648 - 3466 = -818 \text{ kJ mol}^{-1}$$

The negative sign shows that the reaction is exothermic (endothermic reactions have a positive sign).



- 4 Calculate the energy change when hydrogen reacts with oxygen:  
 $2(\text{H}-\text{H}) + \text{O}=\text{O} \rightarrow 2(\text{H}-\text{O}-\text{H})$

Covalent bond	Bond energy ( $\text{kJ mol}^{-1}$ )
C–O	358
C–H	413
H–H	436
O–H	464
O=O	498
C=O	805

C a selection of bond energies

### Did you know?

The strength of a given covalent bond differs between substances, and even between different places in the same molecule, so tables usually show mean bond energies.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** What happens in exothermic and endothermic reactions when bonds are broken and made?
- S2** What information do reaction profiles give us?

### Extend

- E1** Calculate the energy change when methanol undergoes complete combustion:
- $$\text{CH}_3\text{OH} + 1\frac{1}{2}\text{O}_2 \rightarrow \text{CO}_2 + 2\text{H}_2\text{O}$$

### Exam-style question

Explain, in terms of the energy involved in the breaking and making of bonds, why some reactions are endothermic. (2 marks)



## Rates of reaction

Dilute hydrochloric acid reacts with solid calcium carbonate. A student carries out two experiments involving this reaction. She uses the same concentration of hydrochloric acid and the same mass of calcium carbonate each time. The table shows her results.

	Experiment 1	Experiment 2
calcium carbonate	lumps	powder
temperature of acid (°C)	22	45
reaction time (s)	225	25

Evaluate the student's results, explaining any difference in the rate of reaction between the two experiments in terms of particles. (6 marks)

### Student answer

I think the data shows that the reaction in experiment 2 is nine times faster than the same reaction in experiment 1 [1]. The greater rate of reaction in experiment 2 is for two reasons. Firstly, the particles in the powder have a greater surface area [2] than the particles in the lumps used in experiment 1. This means that more particles are exposed to the acid and there are more frequent collisions between reactant particles [3], so the rate of reaction is greater.

The increased rate of reaction is also due to the increased temperature of the acid. The reactant particles in experiment 2 have more energy than those in experiment 1. So, a greater proportion of them have the activation energy or more. This means that the frequency of successful collisions is greater, causing a greater rate of reaction.

It would have been better to vary the size of the particles and the temperature separately, but the data clearly shows that the rate of reaction in experiment 1 is greater than in experiment 2 [4].

[1] The reaction times given in the table are used to make a conclusion about the rates of reaction.

[2] An accurate answer should refer to the surface area to volume ratio of the particles, not just to their surface area.

[3] The answer should refer to successful collisions (as it does later), not just to collisions.

[4] The answer finishes by looking at the data and the experiment, and evaluating how strong the conclusion is.

### Verdict

This is a strong answer. It comes to a clear conclusion about the data. The answer uses scientific knowledge to link the greater rate of reaction in experiment 2 to the energy and frequency of particle collisions.

#### Exam tip

When you are asked to evaluate data, you must use it to form a conclusion. This may include interpreting and explaining the information, or considering its strengths and weaknesses.

## Paper 2

### SC20 Fuels / SC21 Earth and Atmospheric Science

Crude oil and natural gas are mostly used to provide fuels. However, burning oil and natural gas produces carbon dioxide, which is thought to be changing our atmosphere and climate. The photo shows a jet fighter, in which the fuel is kerosene (one of the 'fractions' that crude oil can be split into). The long flame is caused by its afterburner. Extra fuel is injected into the hot exhaust gases, producing additional thrust.

Crude oil also gives us the raw materials needed to make a huge range of products, including polymers such as polythene. Crude oil is, though, a finite resource and the substances we make from it are non-renewable. They will run out one day if we continue to use them.

#### The learning journey

Previously you will have learnt at KS3:

- that mixtures may be separated using fractional distillation
- about fuels and energy resources
- about the acidity of non-metal oxides
- about the production of carbon dioxide by human activity and the impact on climate.

In this unit you will learn:

- about the hydrocarbons found in crude oil and natural gas
- how crude oil is separated into useful fractions
- about the alkanes as an homologous series
- about the problems caused by some atmospheric pollutants
- how and why cracking of oil fractions is carried out
- about the advantages and disadvantages of different fuels for cars
- about how the Earth's atmosphere has changed in the past and how it is changing now
- more about the causes and effects of climate change.





# SC20a Hydrocarbons in crude oil and natural gas

Specification reference: C8.1; C8.2; C8.15

## Progression questions

- What are hydrocarbons?
- Why is crude oil so useful?
- Why is crude oil a non-renewable, finite resource?



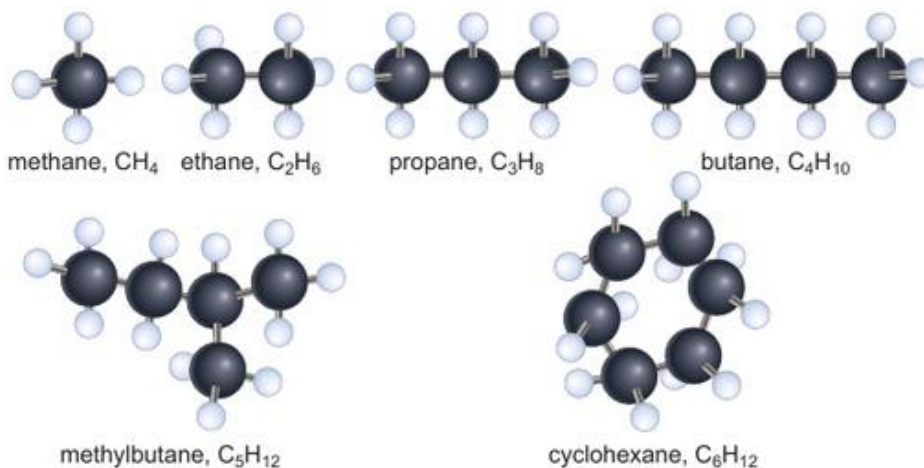
**A** Hydraulic fracturing or 'fracking' is one way to obtain fossil fuels. A mixture of water, sand and other substances is injected into underground rock at high pressure. Natural gas or crude oil flows from cracks in the rocks to the well head at the surface.

**Natural gas** and **crude oil** are natural resources formed from the ancient remains of microscopic animals and plants that once lived in the sea. These remains became covered by layers of sediment. Over millions of years, the remains gradually turned into natural gas and crude oil. The sediments turned into rock, trapping the gas and oil. These resources are **finite resources** because they are not made any more (or are being made extremely slowly), which limits the amounts available to us.

## Crude oil

Crude oil is a complex mixture of **hydrocarbons**. A hydrocarbon is a compound that contains hydrogen and carbon atoms only. Carbon atoms can each form four covalent bonds, so the carbon atoms in hydrocarbon molecules are able to join together in different ways, forming chains and rings. The number of carbon atoms in a hydrocarbon molecule can vary from just one carbon atom to many hundreds of them.

- 5th** 1 Explain why natural gas and crude oil are finite resources.



**B** These ball and stick models show some hydrocarbon molecules. The black spheres represent carbon atoms.

Hydrocarbons exist in different physical states, depending on the size and complexity of their molecules. Crude oil itself is liquid at room temperature, with hydrocarbons in the solid and gas states mixed with hydrocarbons in the liquid state.

- 4th** 2 Explain what is meant by the term hydrocarbon.

- 7th** 3 Explain why carbon atoms can be arranged in chains or rings.

Crude oil is an important source of useful substances, including:

- fuels for vehicles, aircraft, ships, heating and power stations
- **feedstock** or raw materials for the petrochemical industry.

**Petrochemicals** are substances made from crude oil, such as poly(ethene) and other polymers.



4 Describe two reasons why crude oil is useful to us.

## Fuels

**Natural gas** is a mixture of hydrocarbons in the gas state. Methane, the main hydrocarbon in natural gas, is useful for cooking.

Several different liquid fuels can be obtained from crude oil, including petrol and diesel oil for vehicles, and kerosene for aircraft. These **fossil fuels** are **non-renewable** – they are being used up faster than they are being formed. If we carry on using these fuels, they will run out one day.



D Natural gas is used for domestic heating. This is a boiler with the cover taken off.



5 Explain why diesel and methane are *non-renewable* fossil fuels.

## Exam-style question

Octane,  $C_8H_{18}$ , is one of the substances found in petrol.

- Explain why octane is a hydrocarbon. (2 marks)
- State why petrol is non-renewable. (1 mark)



C This oil industry engineer is pouring some crude oil into a container, ready for a test.

## Did you know?

The barrel is the unit of volume used by the oil industry. One barrel of oil is 159 litres. The world uses about 96 million barrels of oil each day – about 180 000 litres per second.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1 What are the main features of crude oil and the substances it contains?
- S2 Why are crude oil and natural gas useful?

## Extend

- E1 Why is crude oil an important finite source of non-renewable substances?



# SC20b Fractional distillation of crude oil

Specification reference: C8.3; C8.4; C8.5

## Progression questions

- How is crude oil separated into useful fractions?
- What are the names and uses of the main fractions from crude oil?
- What are the differences in the molecules found in different fractions from crude oil?



**A** Bitumen, a fraction obtained from crude oil, is mixed with small stones to make road surfaces.

Crude oil is usually not runny enough or ignited easily enough for it to be useful as a fuel. The different hydrocarbons it contains must be separated into simpler, more useful mixtures. This can be achieved using **fractional distillation** because the different hydrocarbons have different boiling points.



- 1** State why crude oil can be separated using fractional distillation.

## In the fractionating column

The industrial fractional distillation of crude oil happens in a tall metal **fractionating column**. Crude oil is heated strongly to **evaporate** it, and the hot vapours are piped into the bottom of the column, where:

- the column is hottest at the bottom and coldest at the top
- the vapours rise through the column and cool down
- the vapours **condense** when they reach a part of the column that is cool enough (below their boiling points)
- the liquid falls into a tray and is piped away
- the vapours with the lowest boiling points do not condense at all and leave at the top as a mixture of gases
- bitumen has the highest boiling point and leaves at the bottom as a hot liquid.

The separated liquids and gases are called oil **fractions** because they are only parts of the original crude oil.



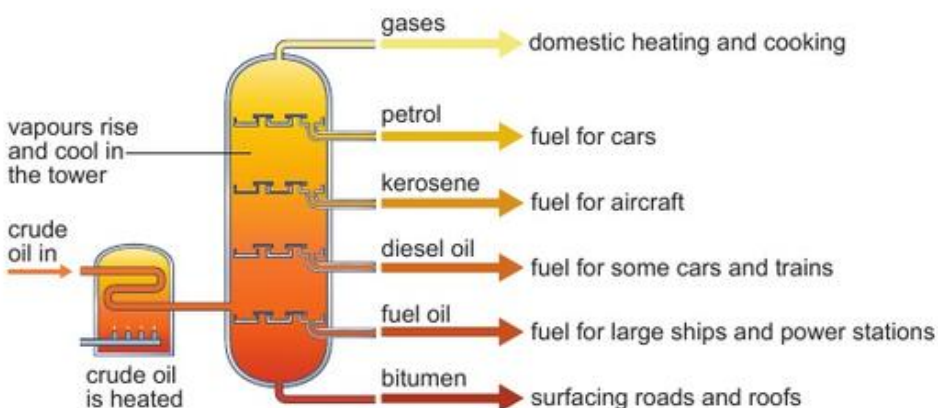
**B** Fractional distillation of crude oil happens at oil refineries.



- 2** Make a table to summarise the names and uses of the six main oil fractions.



- 3** Explain where changes of state happen during fractional distillation of crude oil.



**C** These are the names and uses of the main fractions leaving an oil fractionating column.





## Properties of fractions

Each fraction is a mixture of hydrocarbons, rather than a pure hydrocarbon. However, the hydrocarbons in a given fraction have similar numbers of carbon and hydrogen atoms in their molecules, and similar boiling points.



Different fractions have different uses because they have different properties. For example:

- the hydrocarbons in the gases fraction have the lowest **viscosity** (they flow easily) and are easiest to **ignite** (they are easily set alight), making them suitable for use as fuels
- bitumen is solid at room temperature, and waterproof, making it suitable for surfacing roads and roofs.

The other fractions are liquids at room temperature.

Fraction	Number of atoms in molecules	Boiling point	Ease of ignition	Viscosity
gases	smallest (1–4 carbon atoms) 	lowest (<0 °C) 	easy to ignite 	lowest (flows most easily) 
petrol				
kerosene				
diesel oil				
fuel oil				
bitumen	greatest (>35 carbon atoms)	highest (>350 °C)	difficult to ignite	highest (flows with difficulty)

**D** trends in the properties of the fractions leaving an oil fractionating column

-  **4** Describe the properties of kerosene that make it suitable for use as an aircraft fuel.
-  **5** Describe the relationships between the number of carbon atoms in a hydrocarbon molecule and the physical properties of the hydrocarbon.

### Exam-style question

Petrol and fuel oil are different fractions separated from crude oil. Describe the difference in *one* physical property between these two fractions.

(2 marks)

### Did you know?

Over 100 million tonnes of bitumen is used globally each year. Around 85% is used for road building, and nearly all of it can be recycled when a road wears out and needs repairing.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** How does fractional distillation of crude oil work?
- S2** What do the substances in a crude oil fraction have in common?

### Extend

- E1** How is crude oil made into useful mixtures?
- E2** Describe how the hydrocarbons in different fractions differ from each other.



# SC20c The alkane homologous series

Specification reference: C8.5; C8.6

## Progression questions

- What is the main type of hydrocarbon found in crude oil?
- What are the features of an homologous series of compounds?
- Why do alkanes form an homologous series?



**A** Methane,  $\text{CH}_4$ , is the simplest alkane. It is transported in large amounts as liquefied natural gas (LNG). It is distributed as a gas to homes and businesses through a network of pipelines.

The compounds in crude oil fractions are mostly **alkanes**. Alkanes are hydrocarbons that only have single covalent bonds between the atoms in their molecules. The alkanes form a 'family' or **homologous series** of compounds. There are several different homologous series but the members of a series have these features in common:

- the molecular formulae of neighbouring compounds differ by  $\text{CH}_2$
- they have the same general formula (see the next page)
- they show a gradual variation in physical properties, such as their boiling points
- they have similar chemical properties.



- 1** Name the homologous series of compounds that make up most of crude oil.

## Molecular formulae

The table shows information about the first three alkanes.

Name	Molecular formula	Structural formula
methane	$\text{CH}_4$	<pre>  H     H-C-H       H</pre>
ethane	$\text{C}_2\text{H}_6$	<pre>  H   H         H-C---C-H           H   H</pre>
propane	$\text{C}_3\text{H}_8$	<pre>  H   H   H             H-C---C---C-H               H   H   H</pre>

**B** names and formulae of the first three alkanes

Going from ethane to propane, the number of carbon atoms increases by 1 and the number of hydrogen atoms increases by 2. You can see this in their **molecular formulae**, which show the actual numbers of atoms of each element in the molecules of these compounds. The molecular formula changes by  $\text{CH}_2$  going from  $\text{C}_2\text{H}_6$  to  $\text{C}_3\text{H}_8$ . You might find this easier to see in the **structural formulae** of ethane and propane. It looks as if an extra  $\text{CH}_2$  group of atoms is fitted in between the two  $\text{CH}_3$  groups of atoms.



- 2** Describe the difference between the molecular formulae of methane and ethane.

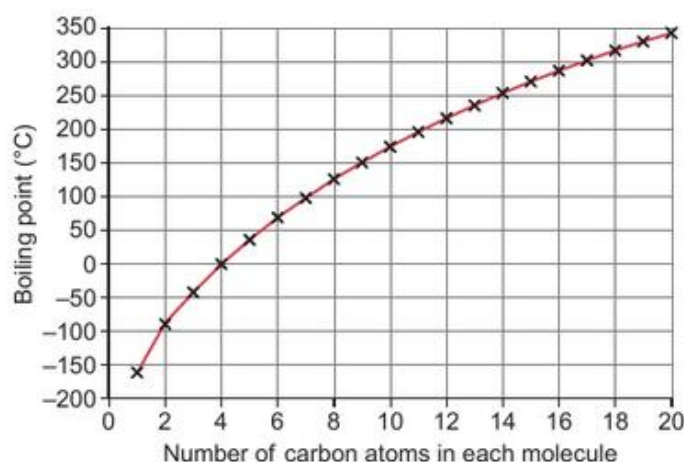
## General formulae

A **general formula** represents the formula for a whole homologous series. The general formula for the alkanes is:  $C_nH_{2n+2}$ . This means that the number of hydrogen atoms is twice the number of carbon atoms, plus two. For example, propane molecules have three carbon atoms, so:

- $n = 3$ , and  $2n+2 = (2 \times 3) + 2 = 8$
- the molecular formula is  $C_3H_8$ .

## Trends in physical properties

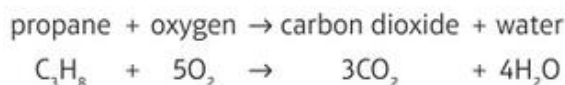
The graph shows how the boiling points of alkanes change as the number of carbon atoms in the molecules increases. Notice that there is a gradual variation in this physical property. The difference in boiling points of the alkanes is the reason why alkanes in crude oil can be separated by fractional distillation.



**C** The boiling point of a straight-chain alkane depends upon the size of its molecules.

## Similar chemical properties

The alkanes have similar chemical properties. For example, they react with excess oxygen to produce carbon dioxide and water. These are the equations for propane, found in camping gas:



**5** Methane reacts with excess oxygen. For this reaction, write:



- the word equation
- the balanced equation.

## Exam-style question

Ethene,  $C_2H_4$ , is a member of the alkene homologous series. The next two alkenes are propene,  $C_3H_6$ , and butene,  $C_4H_8$ . Explain why these three hydrocarbons are members of the same homologous series. (2 marks)

**3** Give the molecular formulae for:



**a** butane (an alkane with four carbon atoms)



**b** hexadecane (an alkane with 16 carbon atoms).

## Did you know?

The carbon atoms in alkanes can be arranged so there are branches as well as straight chains. There are only two ways to arrange the carbon atoms with butane but over 10 000 ways to do this with hexadecane.



**4** Describe how the boiling points of alkanes change as the number of carbon atoms in the molecules changes.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** What are the features of an homologous series?

## Extend

**E1** Explain why the main components of crude oil form an homologous series.

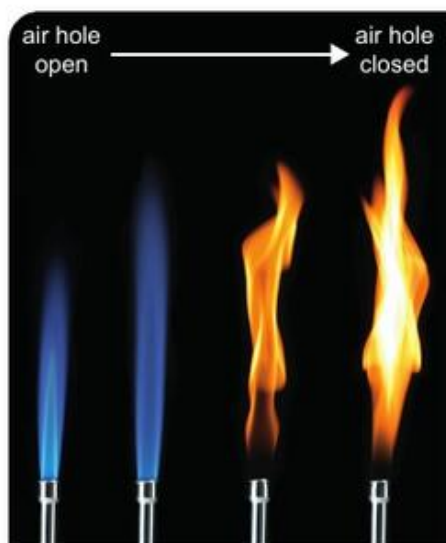


# SC20d Complete and incomplete combustion

Specification reference: C8.7; C8.8; C8.9; C8.10

## Progression questions

- What happens during the complete combustion of a hydrocarbon?
- What happens during the incomplete combustion of a hydrocarbon?
- What problems does incomplete combustion cause?



**A** Adjusting the air hole on a Bunsen burner alters the amount of air reaching the fuel, and changes the flame.

Hydrocarbon fuels react with oxygen in the air when they burn. This is an example of an **oxidation** reaction and is called **combustion**.

## Complete combustion

**Complete combustion** of a hydrocarbon is a reaction in which:

- only carbon dioxide and water are produced
- energy is given out.

Complete combustion happens when there is a plentiful supply of air or oxygen, for example when the air hole on a Bunsen burner is fully open. Methane is the main hydrocarbon found in natural gas:

methane + oxygen → carbon dioxide + water

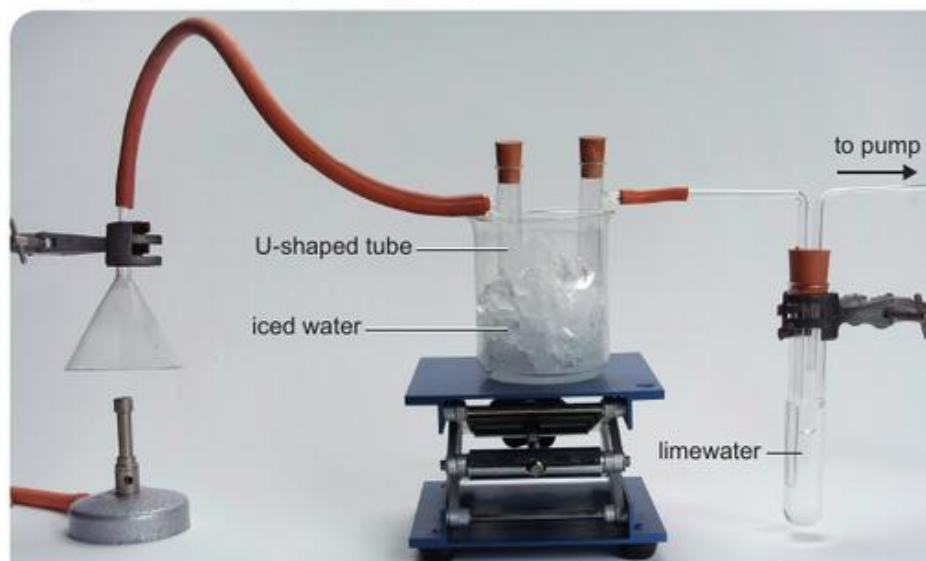


- 1** Write the balanced equation for the complete combustion of pentane,  $\text{C}_5\text{H}_{12}$ .

The apparatus in photo B is used to investigate combustion. A pump draws combustion products from the Bunsen burner through the apparatus. Iced water cools and condenses water vapour passing through the U-shaped tube. White anhydrous copper sulfate in the U-shaped tube turns into blue hydrated copper sulfate, showing the presence of water. Limewater in the boiling tube turns milky, showing the presence of carbon dioxide.



- 2** Describe laboratory tests for water and for carbon dioxide.



**B** This apparatus is used to investigate combustion products.

## Incomplete combustion

**Incomplete combustion** happens when there is a limited supply of air or oxygen, such as when the air hole on a Bunsen burner is closed. During the incomplete combustion of a hydrocarbon:

- water is produced
- energy is given out (but less than with complete combustion)
- **carbon monoxide**, CO, and carbon are produced.

Some carbon atoms in the hydrocarbon may still be fully oxidised to carbon dioxide, but some are only partially oxidised to carbon monoxide. Some carbon atoms are released as smoke and **soot**.

## Incomplete combustion problems

Incomplete combustion can cause problems in appliances that use hydrocarbon fuels, such as boilers and heaters, if they are poorly maintained or unventilated. Carbon monoxide is a **toxic** gas. It combines with **haemoglobin** in **red blood cells**, preventing oxygen combining. This reduces the amount of oxygen carried in the bloodstream, causing affected people to feel sleepy or to become unconscious. Severe carbon monoxide poisoning can even cause death.

Soot can block the pipes carrying away waste gases from an appliance. It blackens buildings, and it can cause breathing problems if it collects in the lungs.



**D** This wall has been blackened by soot from the exhaust pipe of an old boiler.



**5** Describe *one* problem caused by soot.

### Exam-style question

Describe a problem caused by *one* product of the incomplete combustion of natural gas. (2 marks)



**3** Explain why incomplete combustion of hydrocarbons can produce carbon monoxide and carbon.



**C** Carbon monoxide is colourless and odourless, so electronic carbon monoxide detectors are used in homes using natural gas for heating or cooking.



**4** Explain why carbon monoxide is a *toxic* gas.

### Did you know?

After hydrogen, carbon monoxide is the second most common molecule in interstellar space.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Why can burning fuels in a lack of oxygen cause problems?

### Extend

**E1** Explain why adequate ventilation is important when using appliances fuelled by hydrocarbons.



# SC20e Combustible fuels and pollution

Specification reference: C8.11; C8.12; C8.13

## Progression questions

- Why do some hydrocarbon fuels release sulfur dioxide when they are used?
- Why are oxides of nitrogen produced by engines?
- What problems are caused by acid rain?



**A** The trees in this forest have died because of the effects of acid rain.

Carbon dioxide, like other soluble non-metal oxides, forms an acidic solution with water. Rain water is naturally acidic because it contains carbon dioxide from the air. **Acid rain** has a pH lower than 5.2, making it more acidic than natural rain water. Sulfur dioxide is a major cause of this extra acidity.



- 1** Name a substance that causes acid rain.

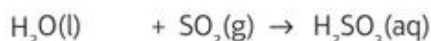


**B** New stonework and similar stonework (on the right) damaged by years of acid rain.

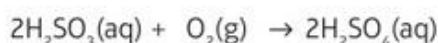
## Sulfur dioxide

Hydrocarbon fuels (such as petrol and diesel oil) may contain sulfur compounds. These occur naturally as **impurities** and are not deliberately added. On the contrary, most of these impurities are removed at oil refineries in an attempt to reduce the environmental problems they cause. When the hydrocarbon fuel is burnt, the sulfur reacts with oxygen to form sulfur dioxide gas,  $\text{SO}_2$ .

Sulfur dioxide dissolves in the water in clouds to form a mixture of acids, including sulfurous acid:



Sulfurous acid is oxidised by oxygen in the air to form sulfuric acid:



This mixture of sulfurous acid and sulfuric acid causes problems when it falls as acid rain. Crops do not grow well when the soil is too acidic. Excess acidity in rivers and lakes prevents fish eggs hatching, and it can kill fish and insects.

Acid rain increases the rate of **weathering** of buildings made of limestone or marble, and breaks down their structures. These rocks are almost pure calcium carbonate, which reacts with sulfuric acid:



Acid rain also increases the rate of corrosion of metals, such as the iron in steel, weakening them:



- 2** Write a balanced equation for the reaction between sulfur and oxygen.



- 3** Describe two environmental problems caused by sulfur dioxide dissolving in rain water.



- 4** Write the balanced equation for the reaction between calcium carbonate,  $\text{CaCO}_3$ , and sulfuric acid.



## Oxides of nitrogen

Car engines are 'internal combustion' engines – fuel is mixed with air and ignited *inside* the engine. This causes temperatures high enough for nitrogen and oxygen in the air inside the engine to react together. The reactions produce various **oxides of nitrogen**, or  $\text{NO}_x$ , which are atmospheric **pollutants**.



**C** Spreading powdered limestone on a field reduces the acidity of the soil.

$\text{NO}_x$  are a cause of acid rain. Nitrogen dioxide,  $\text{NO}_2$ , forms dilute nitric acid when it dissolves in the water in clouds. Nitrogen dioxide is a toxic red-brown gas, and it can cause respiratory diseases such as bronchitis. Catalytic converters in cars convert most of the  $\text{NO}_x$  in exhaust gases to harmless nitrogen.



**D** In sunlight, oxides of nitrogen react with other pollutants to form a harmful 'photochemical smog'.

**6** Catalytic converters can reduce nitrogen dioxide, forming nitrogen and oxygen. Write a balanced equation for this reaction.

### Exam-style question

Large ships use fuel oil, which contains sulfur as an impurity. Explain why this can cause environmental problems when ships travel the oceans. (3 marks)

### Did you know?

Sulfur dioxide causes atmospheric cooling when it is released in volcanic eruptions. Scientists have suggested deliberately injecting sulfur dioxide high into the atmosphere as a way to reduce global warming.

**5** Explain why:



**a** the use of some hydrocarbon fuels causes the production of sulfur dioxide



**b** oxides of nitrogen may be produced when hydrocarbon fuels are used.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Why are some non-metal oxides produced when hydrocarbon fuels are used in cars?
- S2** What problems are caused by acid rain?

### Extend

- E1** How does the use of hydrocarbon fuels produce pollutants?
- E2** Why might the problems caused by acid rain vary over time?



# SC20f Breaking down hydrocarbons

Specification reference: C8.14; C8.16; C8.17

## Progression questions

- Why is cracking needed?
- What happens during the cracking of crude oil fractions?
- What are the advantages and disadvantages of hydrogen and petrol as vehicle fuels?



**A** This storage building has stockpiles outside because the supply of the products is greater than the customers' demand for them.

**1** Use bar chart B to identify:

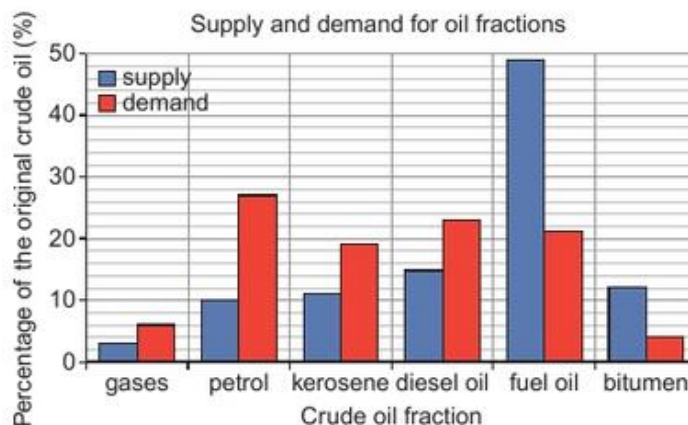


**a** the fractions in higher supply than demand



**b** the fractions in higher demand than supply.

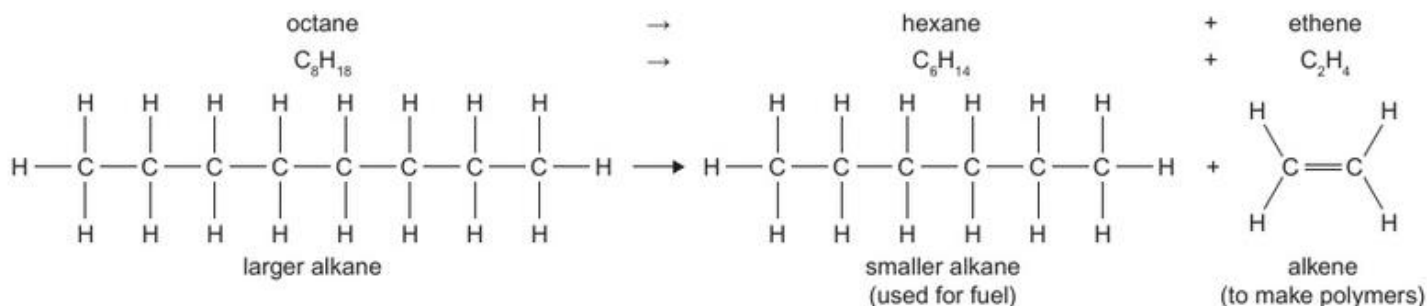
When crude oil is separated by fractional distillation, the volume of each fraction usually does not match the volume that can be sold. The supply of some fractions is greater than customer demand, while the demand for other fractions is greater than the supply. Oil refineries use **cracking** to match supply with demand. Cracking also produces hydrocarbons with C=C bonds. These hydrocarbons are used to make polymers.



**B** example supply and demand for oil fractions

## Cracking

Cracking involves breaking covalent bonds in hydrocarbon molecules. Crude oil fractions are heated to evaporate them. The vapours are passed over a **catalyst** containing aluminium oxide and heated to about 650°C. This speeds up reactions that break down larger hydrocarbon molecules. Smaller, more useful, hydrocarbon molecules form and these can be used as fuels or for making polymers. For example:



**C** a cracking reaction (modelled using structural formulae)

Notice that one of the products, ethene  $C_2H_4$ , is an **alkene**. Alkenes form a different homologous series to alkanes. Alkanes and alkenes are hydrocarbons, but:

- alkanes are **saturated** (their carbon atoms are joined by single bonds,  $C-C$ )
- alkenes are **unsaturated** (they contain a carbon-carbon double bond,  $C=C$ ).

## Fuels for cars

Petrol is in high demand for use as a fuel for cars. It is liquid at room temperature, so large amounts can be stored in the car's fuel tank and then pumped to the engine. Petrol is easily ignited, and its combustion releases large amounts of energy.

Hydrogen can also be used to fuel cars. It is a by-product of cracking and can also be produced by reacting methane (from natural gas) with steam. Unlike petrol and other hydrocarbon fuels, the combustion of hydrogen produces water vapour but no carbon dioxide. Hydrogen fuel has environmental benefits because carbon dioxide is a **greenhouse gas**, linked to global warming and climate change.

Like petrol, hydrogen is easily ignited and its combustion releases large amounts of energy. Unlike petrol, hydrogen is a gas at room temperature. This makes hydrogen difficult to store in large amounts unless it is compressed under high pressure or liquefied by cooling.



**D** In this car, electricity for an electric motor is generated by a fuel cell that uses hydrogen.

- 5** Explain, in terms of hydrogen molecules, an advantage of storing hydrogen in the liquid state rather than in the gas state.

## Exam-style question

Ethene,  $C_2H_4$ , is used to make poly(ethene). Ethene is produced by cracking crude oil fractions.

- a** Explain what is meant by the term cracking. (2 marks)
- b** Complete this equation:  $C_8H_{18} \rightarrow 3C_2H_4 + \dots$  (1 mark)

- 2** Copy and complete this equation, which models a cracking reaction.



- 3** Describe the difference between saturated and unsaturated hydrocarbons.

- 4** Write a balanced equation, with state symbols, for the reaction between hydrogen and oxygen.

## Did you know?

Scientists have discovered that graphene can store hydrogen. Roasted coconut or roasted chicken feathers work too.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** How are crude oil fractions made more useful?
- S2** What are the advantages and disadvantages of using hydrogen in cars, rather than petrol?

## Extend

- E1** How and why are relatively large alkanes broken down?
- E2** What influences the choice of hydrogen or petrol as a fuel for cars?



# SC21a The early atmosphere

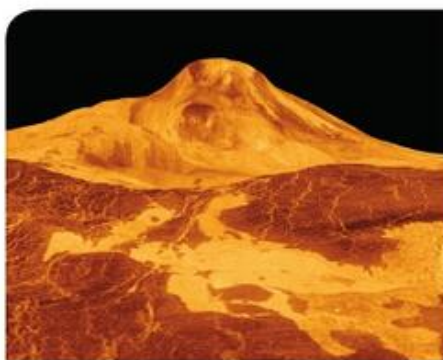
Specification reference: C8.18; C8.19; C8.20

## Progression questions

- What are the names of some common gases produced by volcanic activity?
- What evidence is there for the composition of the Earth's early atmosphere?
- How do scientists explain the formation of the oceans?



**A** Venus' atmosphere is 96.5% carbon dioxide and 3.5% nitrogen.



**B** Maat Mons (an 8 km high volcano on Venus) photographed by the NASA Magellan space probe

Evidence indicates that significant changes have occurred to the **composition** of the Earth's **atmosphere** in its 4.5 billion year history. More recent changes are due to living organisms.

To investigate the atmosphere before life evolved, scientists look at evidence on Earth and study the atmospheres of other planets and moons. These bodies are unlikely to contain life, so their atmospheres may be similar to the Earth's early atmosphere.



- 1** Explain why scientists study the atmospheres of other bodies in our Solar System.

## Earth's early atmosphere

The Earth's early atmosphere is thought to have been mainly carbon dioxide, with smaller amounts of water vapour and other gases, and little or no oxygen.

Volcanoes affect the atmosphere by releasing large amounts of some gases, such as carbon dioxide and water vapour, and small amounts of other gases (including nitrogen). There was a lot of **volcanic activity** on the early Earth, so volcanoes probably helped form its atmosphere.

Earth, Venus and Mars are rocky planets with volcanoes. The atmospheres of Venus and Mars are mainly made of carbon dioxide, thought to have been released by volcanoes. This supports the idea that Earth's early atmosphere also contained lots of carbon dioxide.

Some scientists think that the Earth's early atmosphere was mainly nitrogen. Evidence for this comes from Titan, a moon of Saturn. Titan's atmosphere is 98.4% nitrogen, probably also released by volcanoes. However, space probes have shown that Titan has an icy interior, unlike Earth, Venus or Mars. So, Earth's early atmosphere is less likely to have been like Titan's.



- 2** State how volcanoes affect the atmospheres of planets.



- 3 a** Give the name of the gas most likely to have formed most of the Earth's early atmosphere.



- b** Explain why the discovery of Titan's icy interior does not support the idea that the Earth's early atmosphere was mainly nitrogen.



- 4** Explain why scientists think that the early Earth had an atmosphere similar to Venus.

## Did you know?

The largest volcano in the Solar System is on Mars. Called *Olympus Mons*, it is 27 km high (three times the height of Mount Everest).

## The oceans

About 4 billion years ago, the Earth cooled down. This caused water vapour in the atmosphere to condense to liquid water, which formed the oceans.



- 5 Describe how the Earth's oceans may have formed.

## Oxygen

While the exact composition of the Earth's early atmosphere is uncertain, there is much more direct evidence to support the idea that it contained little or no oxygen.

Oxygen is not produced by volcanoes. Further evidence comes from iron pyrite, a compound that is broken down by oxygen and so only forms if there is no oxygen. It is often found in very ancient rock.

About 2.4 billion years ago, rocks containing bands of iron oxide started to form. This oxidation of iron suggests that oxygen levels increased at this time. There is fossil evidence of microorganisms that may have produced this oxygen. So, scientists think that oxygen from these microorganisms reacted with iron in the early oceans, to produce insoluble iron oxides that formed layers on the seabed.

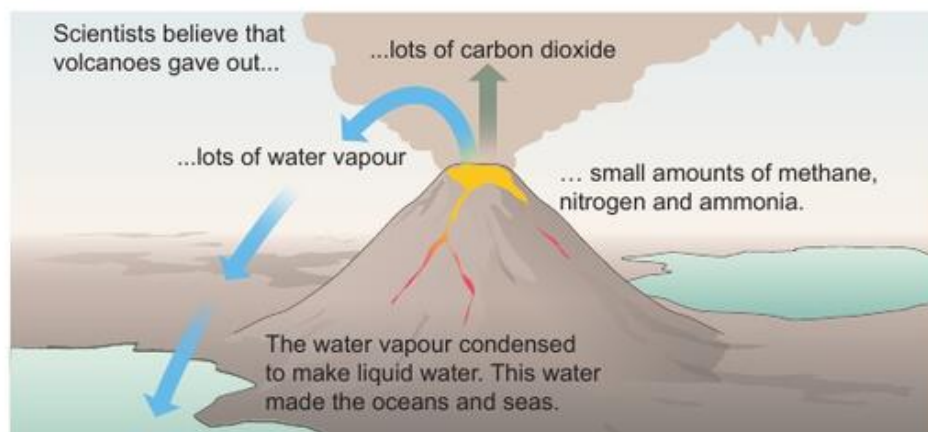
Some geologists suggest that it was only after microorganisms had produced enough oxygen to oxidise the iron in the ocean that atmospheric oxygen levels could rise.



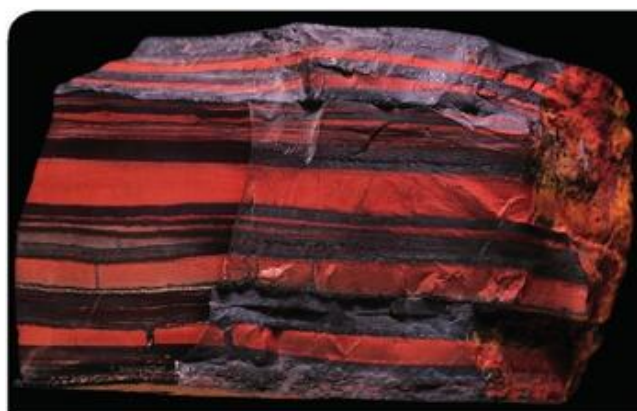
- 6 Explain the evidence that supports the idea that oxygen was not present in the early Earth's atmosphere.



- 7 Why are scientists more certain about the oxygen content of the early atmosphere compared to other gases?



C Many scientists think that the gases in Earth's early atmosphere came from volcanoes.



D bands of iron oxide in an ancient rock

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1 Describe two pieces of evidence that suggest that the composition of the Earth's early atmosphere was mainly carbon dioxide, with little or no oxygen.

### Extend

- E1 Explain why many scientists think that the Earth's early atmosphere was similar to that of Mars today (which is 95.3% carbon dioxide, 2.7% nitrogen, 1.6% argon and 0.13% oxygen).

### Exam-style question

Describe how the Earth's cooling 4 billion years ago caused a change in the composition of the atmosphere. (2 marks)



# SC21b The changing atmosphere

Specification reference: C8.21; C8.22; C8.23

## Progression questions

- Why did the amount of carbon dioxide in the atmosphere change?
- How did primitive organisms change carbon dioxide and oxygen levels?
- What is the test for oxygen?



**A** Shells from dead sea creatures form layers of sediment that turn into sedimentary rocks (e.g. limestone) over millions of years. Shells can often be seen in limestone.



**B** Stromatolites are caused by oxygen-releasing microorganisms called cyanobacteria.

As the young Earth changed, so did the atmosphere. Over hundreds of millions of years the amount of oxygen increased and carbon dioxide levels decreased. Scientists have put forward many **hypotheses** to explain this. There is strong evidence to support some of these.

## Oceans

As the young Earth cooled, water vapour in the atmosphere condensed and formed oceans. Many scientists think that carbon dioxide then dissolved in the oceans, reducing the amount of carbon dioxide in the atmosphere. Sea creatures used the dissolved carbon dioxide to form shells made of calcium carbonate,  $\text{CaCO}_3$ . This then allowed more carbon dioxide to dissolve in the oceans.



- 1** State what happened to the water vapour in the Earth's atmosphere when temperatures cooled.

## Oxygen

Some organisms use energy from the Sun to make food by **photosynthesis**. These organisms change the atmosphere because photosynthesis uses up carbon dioxide and releases oxygen.

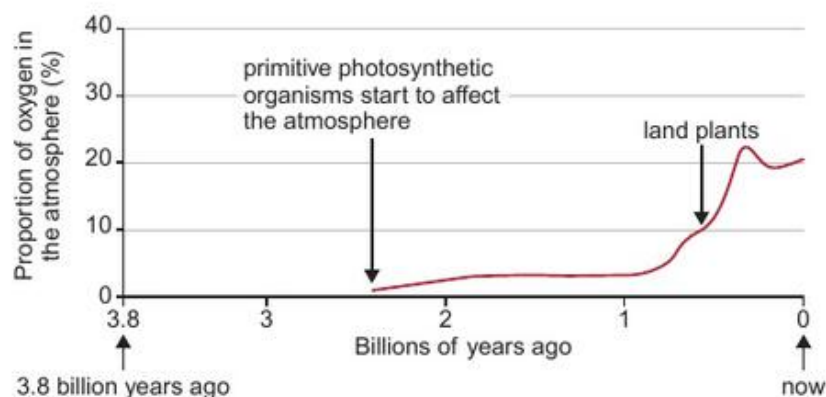
Some of the earliest photosynthetic organisms were cyanobacteria, which live in shallow waters. These bacteria grow in huge colonies and produce sticky mucus. The mucus traps a layer of sand grains and other sediments. The organisms need to move above the sediment layer in order to get sunlight. Over time, the sediment layers build up to form rocky shapes, called stromatolites.



- 2** Explain why cyanobacteria need to move above the sediment as it collects in layers on a stromatolite.

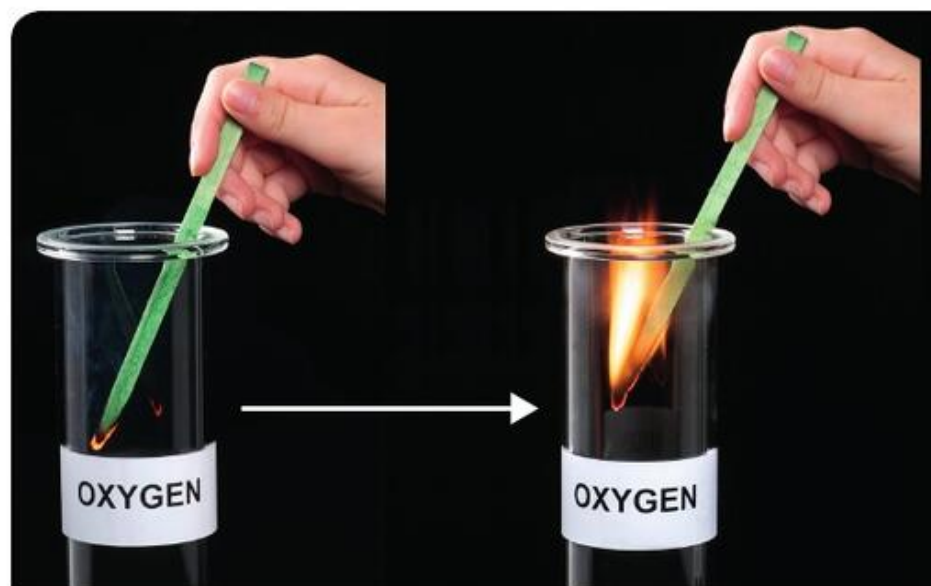
Some stromatolites are over 3 billion years old. They provide evidence that photosynthetic organisms were living at this time. It is thought that microorganisms like these caused a rise in oxygen levels in the oceans and then the atmosphere.

Cyanobacteria evolved into other forms of life, including plants. When land plants evolved, about 500 million years ago, there was another jump in atmospheric oxygen levels.



**C** Diagram showing how some scientists think that oxygen levels in the Earth's atmosphere may have changed over 3.8 billion years.

Today, oxygen makes up about 21% of the Earth's atmosphere. It is important for aerobic respiration in organisms. It also allows combustion (burning) to happen, and this property is used to test for the gas in the laboratory; pure oxygen will relight a glowing splint.



**D** the test for oxygen

- 5** a State the laboratory test for oxygen.
- 6** b Why does oxygen relight a glowing splint?

### Exam-style question

Explain why the formation of oceans on Earth may have caused a decrease in atmospheric carbon dioxide levels.

(3 marks)

- 3** Explain why the presence of ancient stromatolites supports the idea of photosynthesis starting to occur over 3 billion years ago.
- 4** Describe how the evolution of land plants changed the composition of the atmosphere.

### Did you know?

Oxygen is very reactive and would soon disappear from the atmosphere without photosynthetic organisms. Space scientists are very interested in trying to find other planets that contain oxygen in their atmospheres.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Explain how carbon dioxide levels in the Earth's atmosphere decreased.
- S2** Explain how oxygen levels in the Earth's atmosphere increased.

### Extend

- E1** Explain how the development of life on Earth influenced levels of carbon dioxide and oxygen in the atmosphere.

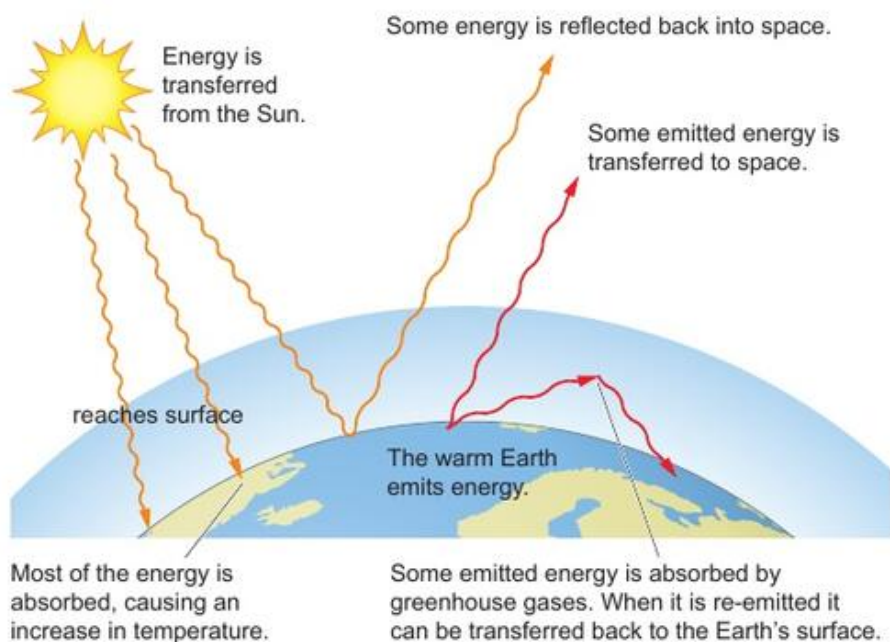


# SC21c The atmosphere today

Specification reference: C8.24; C8.25

## Progression questions

- What are the names of some greenhouse gases?
- How is the greenhouse effect caused?
- What is the link between fossil fuel combustion and climate change?



**A** The greenhouse effect keeps the Earth warm.

Energy from the Sun is transferred to the Earth by waves, such as light and **infrared**. Some energy is absorbed by the Earth's surface, warming it up. The warm Earth **emits** (gives out) infrared waves. Some gases in the air **absorb** energy transferred by these infrared waves. When the gases re-emit the energy, some of it goes back to the Earth's surface and warms it. This is the **greenhouse effect**.

The gases in the atmosphere that absorb energy are called **greenhouse gases**, and include carbon dioxide ( $\text{CO}_2$ ), methane ( $\text{CH}_4$ ) and water vapour ( $\text{H}_2\text{O}$ ). Without them, the mean surface temperature of the Earth would be about  $-18^\circ\text{C}$  (compared to about  $14^\circ\text{C}$ , which it is today).



**1** List three greenhouse gases.

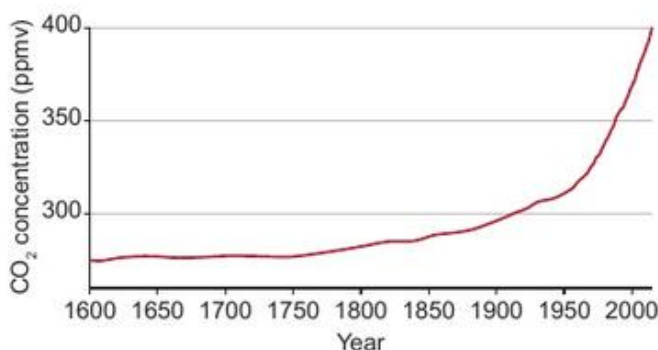


**2** State the influence of the greenhouse effect on the Earth.

## Correlation and climate change

There is evidence to support the idea that human activity is increasing the greenhouse effect and causing **global warming**. This is thought to be causing **climate change** (changes to average weather conditions around the world).

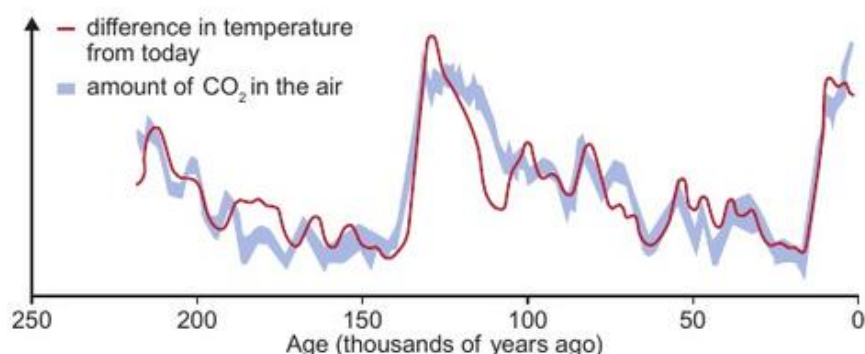
Since about 1850, there has been a steady increase in the burning of fossil fuels for industry. During this period, carbon dioxide levels have increased. We know that combustion releases  $\text{CO}_2$ , and so this is good evidence that increased fossil fuel use has caused increased  $\text{CO}_2$  levels.



**B**  $\text{CO}_2$  levels have risen dramatically since about 1850.

As  $\text{CO}_2$  levels have risen, so has the average temperature of the Earth's surface. There is a strong **correlation** between  $\text{CO}_2$  levels and surface temperature.

However, a correlation does not mean that there is a **causal link** (one thing causes the other). To show a causal link, scientists must collect evidence and explain *how* and *why* the correlation occurs. In this case, scientists want to know whether increasing  $\text{CO}_2$  levels could actually cause global temperatures to rise, whether the reverse happens, or whether there is no causal link.



**C** Average global temperatures and atmospheric carbon dioxide levels are correlated.

Scientists can show in the lab that  $\text{CO}_2$  absorbs infrared. Satellite data confirm that as  $\text{CO}_2$  levels have increased, there has been a reduction in infrared waves from the Earth leaving the atmosphere. This supports the idea that  $\text{CO}_2$  causes temperature rises because it shows how it could occur.

- 3** Why might industrialisation have led to an increase in greenhouse gases?
- 4** **a** Describe what global warming is.  
**b** Compare global warming with climate change.
- 5** **a** What is a 'causal link'?  
**b** Explain why many scientists believe that there is a causal link between recent  $\text{CO}_2$  level increases and a rise in temperature.

## Evaluating the evidence

The amount of carbon dioxide in the air today is measured at monitoring stations around the world. Evidence for historical carbon dioxide levels comes from measuring concentrations of the gas trapped in ice cores. The oldest cores come from Antarctica and give data going back 800 000 years.

The oldest continuous temperature records are for central England and go back to 1659. However, these records cannot be used to assess global temperature changes because they are from only one place. Continuous temperature measurements from around the world exist from about 1880.

Earlier measurements were not very accurate. Modern thermometers are less prone to error and have a greater **resolution**. Today, we can also analyse huge amounts of data from around the world, including temperature measurements from many different sources (such as sensors and satellites).

- 6** How do scientists measure carbon dioxide levels from the past?

## Exam-style question

Describe why the greenhouse effect leads to increased average global temperatures. (3 marks)



**D** Gases are trapped in ice cores.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** Describe the evidence to support the idea that increasing levels of greenhouse gases have caused increased average global temperatures.

## Extend

**E1** Describe how scientists would collect evidence to support a causal link between carbon dioxide levels and global temperatures.



# SC21d Climate change

Specification reference: C8.26

## Progression questions

- What human activities influence the climate?
- What problems might climate change cause?
- How might we limit the impact of predicted climate change?



**A** Methane being released and burnt off on an oil rig.

The increased burning of fossil fuels has released more and more carbon dioxide ( $\text{CO}_2$ ) into the atmosphere, which is thought to be causing global warming. However,  $\text{CO}_2$  is not the only greenhouse gas produced by human activity.

Methane ( $\text{CH}_4$ ) is a more powerful greenhouse gas than  $\text{CO}_2$  because it is much better at absorbing infrared radiation from the Earth. Methane is the main component of natural gas, and is released into the atmosphere when oil and natural gas are extracted from the ground and processed.

Livestock farming (especially cattle) also produces a lot of methane. Cattle have bacteria in their stomachs to digest tough grass. Some of the bacteria produce methane. Soil bacteria in landfill sites and in rice 'paddy' fields also produce a lot of methane.

- 4<sup>th</sup>** 1 State three human activities that cause climate change.
- 6<sup>th</sup>** 2 How does livestock farming increase levels of greenhouse gases in the atmosphere?
- 5<sup>th</sup>** 3 Describe why an increasing world population might lead to more greenhouse gases being released into the atmosphere.



**B** Rice paddy fields produce significant amounts of methane.

## Effects of climate change

Rising average global temperatures will cause ice at the South Pole and glaciers to melt. The extra water will raise sea levels, which will lead to increased flooding in some areas. Higher average temperatures will also result in a loss of 'sea ice' at the poles.

Some animals may move away from their natural habitats to find cooler areas. Some animals and plants may become extinct if they cannot survive at warmer temperatures or find new places to live.

As weather patterns change, some areas will become drier and others will become wetter. Scientists predict that there will be more extreme weather events (such as heavy rainfall, powerful storms and heat waves). These changes will affect wildlife and the growth of crops that people depend upon.

## Did you know?

Sea levels have been rising at a rate of over 3 mm per year since 1993.



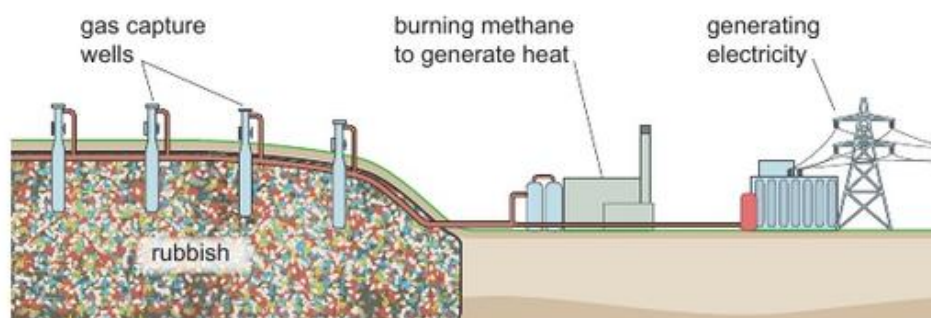
As more  $\text{CO}_2$  is released, more of this acidic gas will dissolve in seawater, lowering its pH. This can harm organisms living in the seas and oceans. Additionally, as ocean temperatures rise, it causes coral to push out the photosynthetic algae that live in their tissues. These algae provide the colour of coral and so coral 'bleaching' may occur.

- 4** List four negative effects of climate change.

## Limiting the impact

Using renewable energy resources can reduce greenhouse gas emissions, but there is a risk that this may not be enough to mitigate (lessen) the effects of climate change that we are already seeing.

Some people have suggested global engineering solutions, to reflect sunlight back into space or to capture  $\text{CO}_2$  from the air and bury it underground. However, all countries will need to work together to reduce emissions and help pay for large-scale engineering. There is a risk that some countries will not help and that delicate ecosystems may be disrupted.



**D** Methane released from landfill sites can be captured and burnt to generate electricity.

Other ways of limiting the impact of climate change involve helping local people to adapt to new conditions. These include building flood defences, dams and irrigation systems. However, these ideas may destroy important habitats and there is a risk that they will not work.

- 5** Explain why an increased use of renewable energy might help to limit the impact of climate change.
- 6** Explain why international cooperation is important in dealing with climate change.
- 7 a** Identify a way of limiting the impact of climate change.
- b** State one risk associated with your suggestion.

## Exam-style question

- a** State a problem caused by climate change. (1 mark)
- b** Describe what can be done to limit any harmful impact. (1 mark)



**C** If coral remain 'bleached' for too long they can die.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** State an effect on the climate of increased carbon dioxide and methane levels, and identify whether the effect is global or local.
- S2** Identify one way of reducing the harmful effect of your example above.
- S3** State a risk associated with your suggested mitigation method.

## Extend

- E1** State the three most important policies you think the government should have for tackling climate change. Justify your choices.

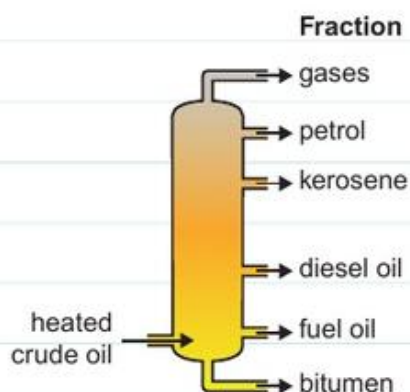


## Fuels

Useful substances are separated from crude oil by fractional distillation. The diagram shows the column used and the fractions it produces. The bitumen fraction leaves from the bottom of this column.

Look at the heights at which the kerosene and fuel oil fractions are collected. Explain the properties and uses for kerosene and fuel oil using the information.

(6 marks)



### Student answer

The top of the fractionating column is cooler than the bottom. Kerosene vapours condense higher up the column than fuel oil. Kerosene has a lower boiling point than fuel oil. This is because the hydrocarbon molecules in kerosene have fewer carbon atoms and hydrogen atoms than the hydrocarbon molecules in fuel oil [1]. Kerosene is more easily ignited than fuel oil [2]. It also has a lower viscosity than fuel oil, so kerosene flows more easily [3]. The differences in these properties make kerosene useful as a fuel for aircraft, [4] while fuel oil is used as a fuel for large ships and some power stations [5].

[1] The answer uses information given in the diagram, and the student's own knowledge and understanding, to link the size of the molecules in each fraction to its boiling point and position.

[2] This describes an important difference in the properties of the two fractions.

[3] It makes clear what low viscosity means – a less viscous substance flows more easily than a more viscous substance.

[4] It is important not to just describe kerosene as used in aircraft, but as a fuel for aircraft.

[5] More than one use of fuel oil is given, and both are correct.

### Verdict

This is a strong answer. It shows good knowledge and understanding of the fractions obtained from crude oil, and uses correct scientific language. The answer uses scientific knowledge to link the size of the hydrocarbon molecules to their boiling points, and therefore the heights at which they are collected in the fractionating column. The answer is organised logically, explaining the differences in properties and then the difference in uses.

### Exam tip

If you are given information in a question, make sure you use and refer to it in your answer. In this case, the question stated that you should use the information from the diagram.

## Paper 2

### **SC22** Hydrocarbons / **SC23** Alcohols and Carboxylic Acids / **SC24** Polymers

Your body contains natural polymers including proteins, starch and DNA. Chemists discovered how to make poly(ethene), an artificial polymer, by accident over a hundred years ago. Nowadays, chemists know a lot more about polymers and how to make them using hydrocarbons, alcohols and carboxylic acids. They are able to develop polymers with astonishing properties, including light-emitting polymers that form flexible screens and polymers that mimic the appearance and flexibility of human skin. This robot head is covered with a spongy polymer, which helps the robot produce different life-like expressions.

#### The learning journey

Previously you will have learnt at KS3:

- about combustion of fuels
- about the properties of polymers.

In this unit you will learn:

- about the structures and properties of alkanes and alkenes
- how a concentrated solution of ethanol is produced from carbohydrates
- about the structures of alcohols and carboxylic acids
- about the chemical properties and uses of alcohols and carboxylic acids
- about the composition of biological polymers
- how poly(ethene) and other polymers are made
- about the disposal and recycling of polymers.





# SC22a Alkanes and alkenes

Specification reference: C9.10C; C9.11C; C9.12C; C9.13C

## Progression questions

- What are the names, formulae and structures of the four smallest alkanes?
- What functional group is present in all alkenes?
- How is the position of this functional group shown in alkene names?



**A** Carpenter ants release the alkane undecane ( $C_{11}H_{24}$ ) to signal danger, which other carpenter ants detect using their antennae.

## Alkanes

**Hydrocarbons** are molecular compounds that contain only carbon and hydrogen. **Alkanes** are a group of hydrocarbon molecules based on a chain of carbon atoms. They all have the **general formula**  $C_nH_{2n+2}$  (where 'n' is a whole number). A series of molecules that have a general formula and vary in a single feature (such as carbon chain length) form an **homologous series**.

- 6<sup>th</sup>** 1 State why alkanes are hydrocarbons.
- 7<sup>th</sup>** 2 An alkane has six carbon atoms. Calculate the number of hydrogen atoms it has.
- 7<sup>th</sup>** 3 In what feature do members of the alkane homologous series vary?

The International Union of Pure and Applied Chemistry (IUPAC) developed the naming system for hydrocarbons. A prefix shows the length of the carbon chain and a suffix shows the homologous series. The suffix for alkanes is '-ane', as shown in table B.

Number of carbons in the chain	Prefix	Alkane	Molecular formula	Structural formula
1	meth-	methane	$CH_4$	<pre>  H     H-C-H       H</pre>
2	eth-	ethane	$C_2H_6$	<pre>  H H       H-C-C-H         H H</pre>
3	prop-	propane	$C_3H_8$	<pre>  H H H         H-C-C-C-H           H H H</pre>
4	but-	butane	$C_4H_{10}$	<pre>  H H H H           H-C-C-C-C-H             H H H H</pre>



**B** the first four members of the alkane homologous series

In table B the lines in the structural formulae represent covalent bonds between atoms. C-H shows a single covalent bond between carbon and hydrogen atoms. C-C shows a single covalent bond between carbon atoms. Molecules are **saturated** if all the carbon-carbon covalent bonds are single bonds. Alkanes are saturated hydrocarbons.

- 6<sup>th</sup>** 4 How many covalent bonds are there in one ethane molecule?
- 6<sup>th</sup>** 5 In an alkane, how many covalent bonds are found attached to every:
  - 6<sup>th</sup>** a carbon atom
  - 6<sup>th</sup>** b hydrogen atom?

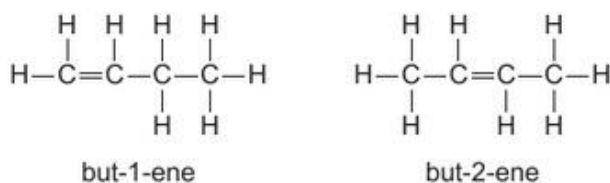
## Alkenes

Hydrocarbons with at least one double covalent bond between adjacent carbon atoms are **unsaturated**. Alkenes are an homologous series of unsaturated hydrocarbons with the general formula  $C_nH_{2n}$ . The first three alkenes in the series are shown in table C. The carbon-carbon double bond ( $C=C$ ) is known as a **functional group**. A functional group is an atom or group of atoms that is mainly responsible for a molecule's chemical properties.

-  **6** What feature is common to all alkenes but not alkanes?
-  **7** How many pairs of electrons are there in a double covalent bond?




## Isomers



A functional group can be in different places in molecules with the same molecular formulae. Molecules with the same molecular formula but different arrangements of atoms are called **isomers**. Butene has two isomers. Numbers are added to the names to show where the  $C=C$  functional group is, as shown in diagram D.



**D** The numbers in the butene isomer names show the position of the double bond. The 1 shows that the  $C=C$  bond starts at the end of the molecule on the first carbon, the 2 shows that it is in the middle. Carbon atoms are numbered starting from the end closest to the double bond.

- 8** In butene, how many of the following bonds are there?

-  **a**  $C-C$
-  **b**  $C-H$
-  **c**  $C=C$

-  **9** Which butene isomer is shown in table C?
-  **10** Suggest why there is not a molecule called but-3-ene.

Name	Molecular formula	Structural formula
ethene	$C_2H_4$	$  \begin{array}{cc}  H & H \\    &   \\  C & =C \\    &   \\  H & H  \end{array}  $
propene	$C_3H_6$	$  \begin{array}{ccccc}  & H & H & H & \\  &   &   &   & \\  H & -C & -C & =C & \\  &   & &   & \\  & H & & H &   \end{array}  $
butene	$C_4H_8$	$  \begin{array}{ccccccc}  & H & H & H & H & & \\  &   &   &   &   & & \\  H & -C & -C & -C & =C & \\  &   &   & &   & & \\  & H & H & & H & &   \end{array}  $

**C** the first three members of the alkene homologous series

### Did you know?

Many plants produce ethene to help ripen their fruits.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Represent one saturated and one unsaturated hydrocarbon using their molecular and structural formulae, together with their names.
- S2** Label your answer to **S1** with the similarities and differences between the two molecules.

### Extend

- E1** Represent a five-carbon alkene using structural and molecular formulae.
- E2** Suggest ways in which isomers of this hydrocarbon may be different.

### Exam-style question

Explain the difference between a saturated and an unsaturated hydrocarbon.

(2 marks)

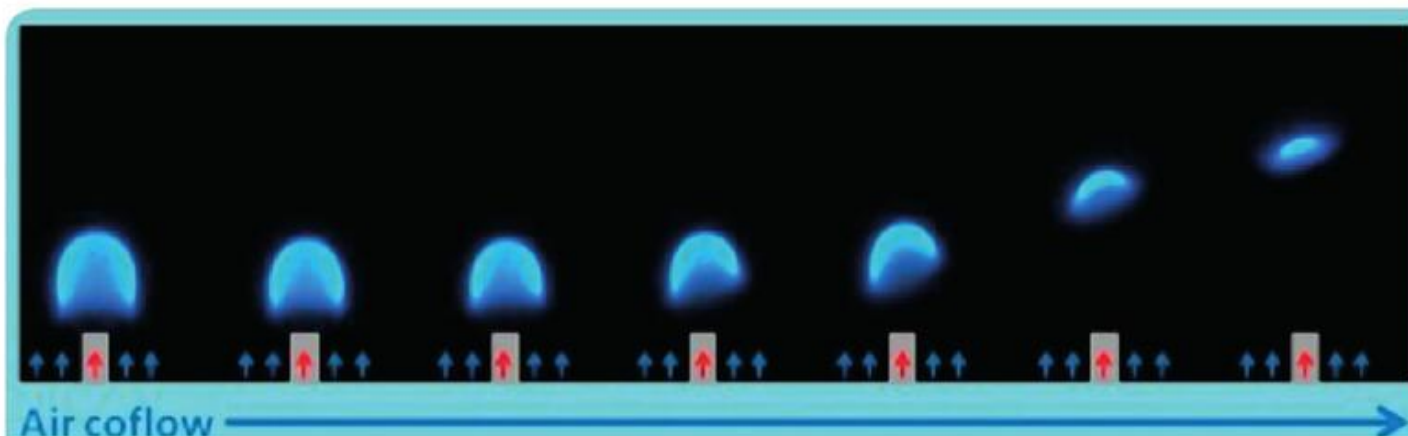


# SC22b Reactions of alkanes and alkenes

Specification reference: C9.14C; C9.15C; C9.16C

## Progression questions

- What products are formed by the complete combustion of hydrocarbons?
- How can bromine water be used to distinguish between alkanes and alkenes?
- What are the structures of the reactants and products when bromine and ethene react?

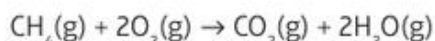


A This is ethene burning in space, where flames are a different shape from those on Earth.

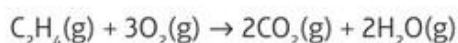
## Combustion

When hydrocarbons burn, carbon dioxide and water are formed when there is a plentiful supply of oxygen. During combustion, **oxidation** occurs. If all the atoms from the hydrocarbon are fully oxidised, the products are only water and carbon dioxide. This is called **complete combustion**.

Methane is the main alkane in natural gas. When it is burnt in a Bunsen burner with the air hole open, the following reaction occurs:



Ethene is the shortest alkene. The complete combustion of ethene is shown in the equation below:



If there is an insufficient amount of oxygen, full oxidation does not occur and carbon (soot) and carbon monoxide can be formed as well. In photo B, the orange flame is due to the presence of soot formed by **incomplete combustion**.



B Fire breathers use a fuel made of alkanes. The orange flame is caused by glowing particles of unburnt soot.

- 1 What are the products of the complete combustion of propene in oxygen?
- 2 What is the functional group found in alkenes?
- 3 Explain why oxidation occurs in these combustion reactions.
- 4 Why do the products of complete combustion not allow a scientist to distinguish easily between alkenes and alkanes?

## The bromine water test

Bromine water is a dilute solution of bromine in water,  $\text{Br}_2(\text{aq})$ . It has an orange-brown colour. When it is mixed with alkenes a chemical reaction occurs, leading to colourless products. Alkanes do not cause decolourisation and so bromine water is used to test between alkenes and alkanes.

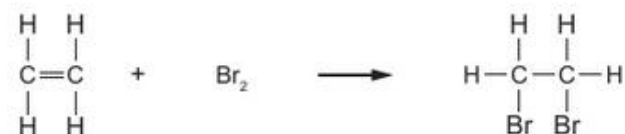
The test works because the  $\text{C}=\text{C}$  double bond reacts with the bromine to form a colourless product. The bromine is therefore removed from the solution, which loses its colour.

- 5** a Describe the effects of bromine water on alkanes and alkenes.  
 b Explain the effects you have described.
- 6** Ripening bananas produce a gas. Describe how you would find out if this gas is an alkene or an alkane.

## Addition reactions

Bromine gas is an orange-brown colour and reacts with alkenes. For example, it reacts with ethene to produce a colourless liquid called 1,2-dibromoethane. The 'di' in the name of the product means 'two' and the numbers show that the two bromine atoms are attached to different carbon atoms. If they were attached to the same carbon atom it would be called 1,1-dibromoethane.

ethene + bromine  $\longrightarrow$  1,2-dibromoethane



**D** Ethene reacts with bromine to form 1,2-dibromoethane.

Diagram D shows that in this reaction two reactant molecules add to one another to form just one product molecule. A reaction in which reactants combine to form one larger product molecule and no other products is called an **addition reaction**.

- 7** Explain what you would see happening during the reaction between ethene and bromine gas.
- 8** Propene reacts with bromine to form 1,2-dibromopropane.
- a Explain why this is an addition reaction.
- b Draw the structure of a molecule of 1,2-dibromopropane, showing all the covalent bonds.

## Exam-style question

Bottles of two liquids, hexane and hexene, have been muddled. Explain how bromine water might be used to distinguish between them. (2 marks)



**C** The effects of bromine water on unsaturated hydrocarbons (containing  $\text{C}=\text{C}$ ) and saturated hydrocarbons (containing  $\text{C}-\text{C}$  only) are different.

## Did you know?

Alkanes used to be called 'paraffins'. This word comes from the Latin 'parum' for 'little' and 'affinis' for 'affinity'. Alkanes do not undergo many chemical reactions and so they have 'little affinity' for other substances.

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** Using the example of ethene reacting with bromine, describe what an addition reaction is.
- S2** Explain why ethane does not undergo this reaction.

## Extend

- E1** But-2-ene reacts with bromine. Draw out the equation for this reaction using structural formulae, and name the product.



# SC23a Ethanol production

Specification reference: C9.33C; C9.34C

## Progression questions

- How are alcoholic drinks produced?
- What chemical reaction occurs during fermentation?
- How can we make alcohol solutions more concentrated?



**A** Drinks that contain a high percentage of ethanol are called spirits.

## Did you know?

The ethanol content of spirits used to be measured in 'degrees proof', where 100° equalled 50% alcohol.



**C** During fermentation by yeast, sugars (such as glucose) are broken down.

Alcoholic drinks have been made for thousands of years. These drinks contain a chemical substance called ethanol (although people often call it 'alcohol'). Its formula is  $C_2H_5OH$ . Ethanol can also be used as a fuel for vehicles and as a raw material for the chemical industry.

- 1 What is the proper chemical name and molecular formula for alcohol?
- 2 Alcoholic drinks can be described as spirits or non-spirits.
  - a What is the difference between them?
  - b Name an example of each type.

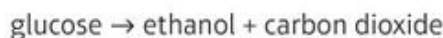
The ethanol in alcoholic drinks is made from **sugars**. Sugars are small, soluble substances that belong to a group called **carbohydrates** (compounds made of carbon, hydrogen and oxygen). Many fruits (such as grapes) contain a lot of sugars.

Seeds contain a carbohydrate called **starch**, which is a long polymer. Starch must be broken down into sugars in order to make alcoholic drinks. Seeds are germinated before using them to make ethanol because, during this process, **enzymes** in the seeds naturally turn the starch into sugars.

Drink	beer	wine	whisky	vodka
Commonly used plant	barley seeds	grapes	barley seeds	wheat seeds

## B plant sources of alcoholic drinks

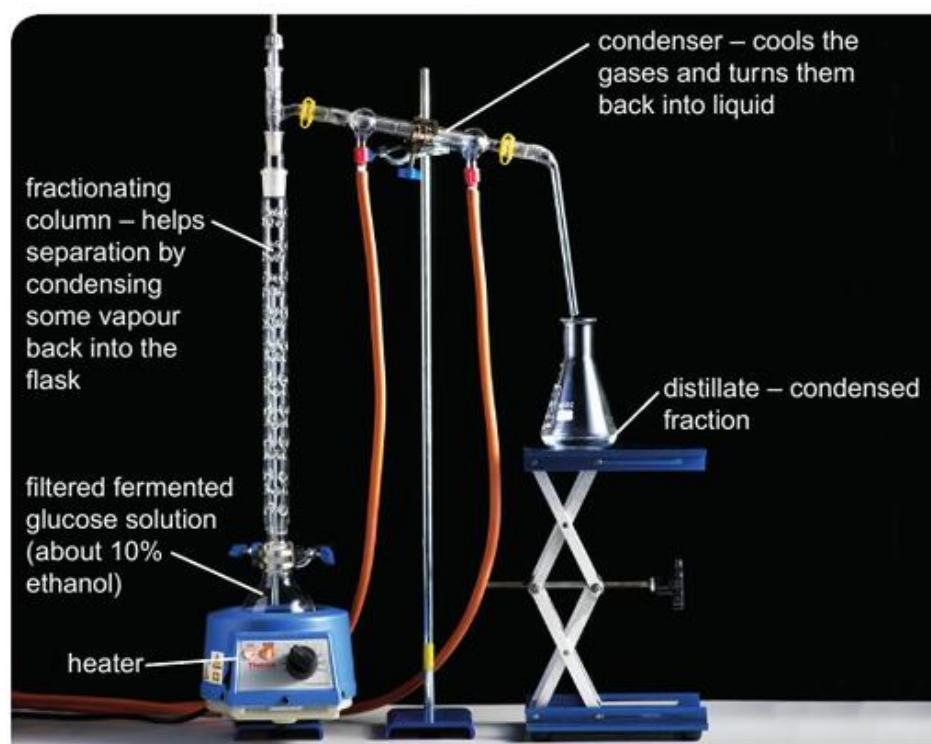
Plant material containing sugars is mixed with water and yeast. Enzymes in the yeast turn the sugars into ethanol and carbon dioxide, in a process called **fermentation**:



- 3 a Where are carbohydrates formed and what elements do they contain?
  - a Give the name of the group of soluble carbohydrates needed for fermentation.

During fermentation, the temperature and pH must be carefully controlled to help the enzymes work at their best. Photo C shows a fermentation flask. The air lock in the neck of the jar allows carbon dioxide to escape while keeping air out. This is necessary because yeast fermentation is a type of **anaerobic respiration** and only occurs in the absence of oxygen.

The fermentation process only produces alcohol concentrations up to 15%, as higher concentrations kill the yeast cells. More concentrated solutions of ethanol are formed by **fractional distillation**, as shown in photo D. This works because the boiling point of ethanol (78°C) is lower than the boiling point of water (100°C). The heated liquids evaporate and their vapours cool as they rise up the fractionating column. As the ethanol has a lower boiling point, it remains as a gas for longer, and separates from the water. As a result, the first **fraction**, or **distillate**, that is collected contains a higher percentage of ethanol.



D fractional distillation of an ethanol solution

- 7** Explain how fractional distillation can be used to increase the concentration of an alcohol solution. Include the following words in your explanation:

boiling point; concentrated; condense; dilute; ethanol; evaporate; heated; water

### Exam-style question

Burning glucose completely in oxygen produces carbon dioxide and water. Compare and contrast the combustion and fermentation of glucose.

(2 marks)

- 4** Describe what is meant by 'anaerobic conditions'.
- 5** Explain why a grape and water mixture must be kept at a certain temperature when making wine.
- 6** Explain why beers rarely have an alcohol concentration greater than 10%.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Draw a flow chart to show the stages involved in the production of a highly concentrated alcohol solution from a source of plant carbohydrates.

### Extend

- E1** In industry, ethanol can be produced by fermentation using a continuous process, in which the reaction is constantly kept going. Write a balanced equation for the fermentation of glucose ( $C_6H_{12}O_6$ ), and explain what would need to be done to keep this reaction going at a constant rate in a continuous fermentation system.

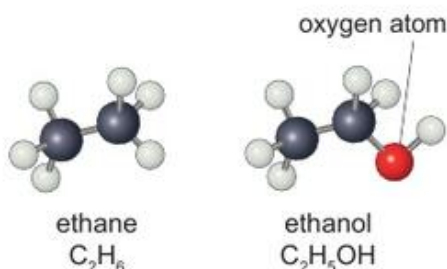


# SC23b Alcohols

Specification reference: C9.26C; C9.27C; C9.32C

## Progression questions

- What are the names, formulae and structures of the four smallest alcohols?
- What functional group is present in all alcohols?
- What are some chemical properties of alcohols?



**A** Ethane and ethanol are both organic compounds but are in different homologous series.

- 1** Explain what is meant by an homologous series.
- 2** Describe one difference and one similarity in the structures of ethane and ethanol.

### Did you know?

Ethanol, propanol and butanol are used in perfumes for two main reasons. They are all good solvents and readily dissolve the ingredients. They also evaporate easily, which helps to spread the perfume's aroma.

**Organic compounds** have a central framework of carbon atoms, onto which hydrogen and other atoms are attached. Ethanol, for example, is organic and has a central framework of two carbon atoms joined by a covalent bond.

Ethanol differs from a similar organic compound called ethane by having an additional oxygen atom. Ethanol and ethane have very different properties and are each in a different family or **homologous series** of compounds. Ethane belongs to the **alkane** series and ethanol belongs to the **alcohol** series.

Compounds in the same homologous series have the same general formula, have similar chemical reactions and display a trend in properties. Their names are also similar. All alcohol names end in '-anol' and have the general formula  $C_nH_{2n+1}OH$  (where  $n$  stands for a number).

Name	Molecular formula	Structural formula
methanol	$CH_3OH$	<pre>  H     H-C-O-H       H</pre>
ethanol	$C_2H_5OH$	<pre>  H H       H-C-C-O-H         H H</pre>
propanol	$C_3H_7OH$	<pre>  H H H         H-C-C-C-O-H           H H H</pre>
butanol	$C_4H_9OH$	<pre>  H H H H           H-C-C-C-C-O-H             H H H H</pre>

**B** The naming of compounds uses a set of rules produced by the International Union of Pure and Applied Chemistry (IUPAC).

All alcohols have similar chemical properties. They all:

- produce carbon dioxide and water on complete combustion, for example:  
 $C_2H_5OH + 3O_2 \rightarrow 2CO_2 + 3H_2O$   
ethanol + oxygen  $\rightarrow$  carbon dioxide + water
- can be oxidised to form compounds called carboxylic acids (see SC23c)
- react with reactive metals, such as sodium, forming hydrogen gas as one of the products.

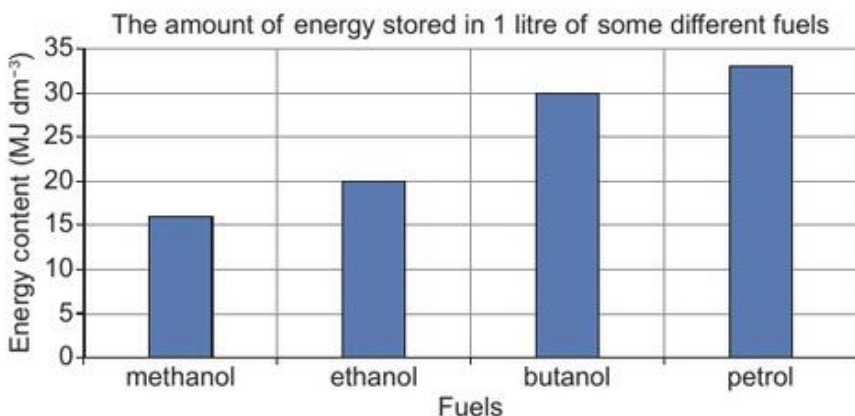
- 3** Describe the difference in molecular structure between propanol and butanol.
- 4** Hexanol has six carbon atoms. How many hydrogen atoms does hexanol have?



**C** Alcohols react with sodium metal but their reactivity depends on carbon chain length.

The atom or group of atoms in a molecule that is responsible for its main chemical reactions is called a **functional group**. Alcohols have similar chemical properties because they all contain one  $\text{-OH}$  functional group.

Alcohols have some important uses. Many alcohols are used as solvents for cosmetics, medical drugs and varnishes. Methanol and ethanol are widely used as fuels and can be made from **renewable sources**. Scientists are now working on developing processes to make propanol and butanol fuels from renewable sources, such as by fermenting plant cellulose using certain bacteria.



**D** Butanol has some advantages over some other alcohols used as fuels.

- 9** State three uses of alcohols.
- 10 a** Explain why butanol produced by fermentation will be a renewable fuel.
- b** Use chart D to suggest why scientists are interested in using butanol as a fuel.

### Exam-style question

The third member of the alcohol series has the molecular formula  $\text{C}_3\text{H}_7\text{OH}$ .

- a** Name this alcohol. (1 mark)
- b** Describe the functional group in this molecule. (2 marks)

**5** Pentanol is an alcohol with five carbon atoms.

- a** State its molecular formula.
- b** Draw a structural formula for pentanol and circle the functional group.
- 6** Describe three chemical properties of pentanol.
- 7 a** Write a word equation for the combustion of butanol.
- b** Write a balanced equation for the combustion of butanol.
- 8** What trend in reactivity is shown in diagram C?

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Construct a table showing the names, formulae, structural formulae and functional groups of the four simplest alcohols.

### Extend

**E1** Explain, by referring to structures and properties, why the alcohols are an homologous series.



# SC23b Core practical – The combustion of alcohols

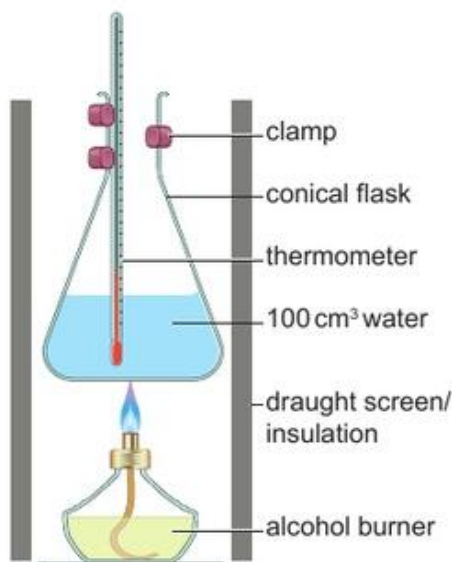
Specification reference: C9.28C

## Aim

Investigate the temperature rise produced in a known mass of water by the combustion of the alcohols ethanol, propanol, butanol and pentanol.



**A** Most modern car engines can run on a fuel that is 90% petrol and 10% ethanol (often called E10). Some cars can run on fuels containing much higher proportions of ethanol, such as E85.



**B** investigating energy in fuels

Traditional fossil fuel energy resources are running out, and their use is also considered to be damaging our environment. We therefore need to find new sources of 'clean' energy. For example, ethanol, formed by the fermentation of sugars, is added to petrol to make it 'greener'. Scientists are now investigating ways of making engines that use other alcohols that are, at present, more difficult to obtain from renewable sources. There are a number of different characteristics that have to be considered when comparing fuels. These include availability, cost, toxicity, ease of combustion, environmental effects and the amount of energy released per unit mass.

## Your task

You will investigate and compare the energy given out by the combustion of different alcohols. To do this, you will measure the temperature rise of a known mass of water caused by the combustion of ethanol, propanol, butanol and pentanol.

## Method

Wear eye protection. Do not refill the alcohol burner if there are any naked flames nearby.

- Measure the mass of an alcohol burner and cap. Record the mass and the name of the alcohol.
- Place the alcohol burner in the centre of a heat-resistant mat.
- Use a measuring cylinder to add 100 cm<sup>3</sup> of cold water to a conical flask.
- Measure and record the initial temperature of the water and clamp the flask above the alcohol burner.
- Light the wick of the burner and allow the water to heat up by about 40 °C.
- Replace the cap on the burner and measure and record the final temperature of the water.
- Measure the mass of the alcohol burner and cap again and record the mass.
- Calculate the mass of the alcohol burned to produce a 1 °C rise in temperature.
- Repeat steps A to H using fresh, cold water and a different alcohol.

## Exam-style questions

- 1 Look at the list of steps in the method described on the previous page.
  - a List six pieces of apparatus required for this investigation. (3 marks)
  - b State the dependent and independent variables. (2 marks)
  - c Explain why fresh water has to be used when repeating the experiment. (2 marks)
  - d State the measurements made during the investigation. (4 marks)
- 2 Look at the investigation apparatus set up in diagram B.
  - a Describe how the position of the flask and the alcohol burner will need to be controlled during each repeat. (1 mark)
  - b Explain the purpose of the draught screen/insulation. (1 mark)
  - c Describe another safety precaution that will need to be taken, in addition to wearing eye protection. (1 mark)
- 3
  - a State four possible sources of error in the investigation described on the previous page. (4 marks)
  - b State the main source of error in this investigation. (1 mark)
- 4 The results obtained by investigating two alcohols are shown in table C.
  - a Calculate the mass of alcohol burned and the temperature rise produced for each of the alcohols. (4 marks)
  - b By analysis of the results in table C, which of the two alcohols releases more energy for each gram burned? (3 marks)

Alcohol	ethanol	propanol
Initial mass of burner + cap (g)	291.60	284.25
Final mass of burner + cap (g)	290.50	285.25
Initial temperature of 100 cm <sup>3</sup> water (°C)	20	20
Final temperature of 100 cm <sup>3</sup> water (°C)	59	62

C

- 5 In another experiment, 1.5 g of butanol produces a temperature rise of 50 °C in 100 cm<sup>3</sup> of water. Calculate the mass of butanol that needs to be burned to cause a 1 °C rise in the same volume of water. (1 mark)

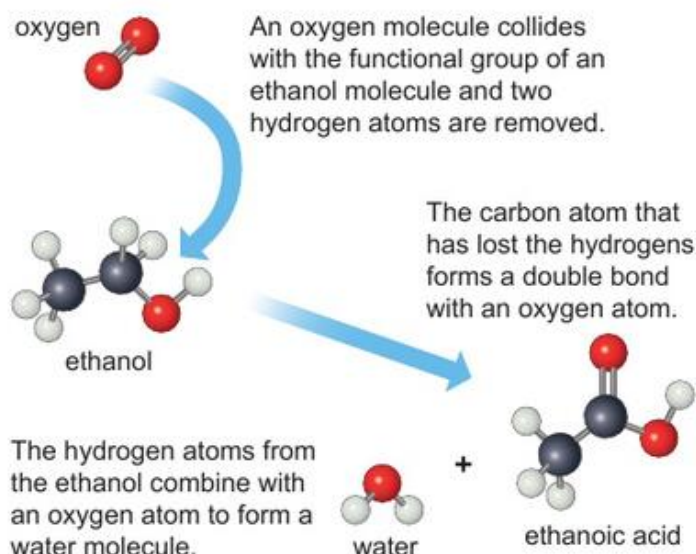


# SC23c Carboxylic acids

Specification reference: C9.29C; C9.30C; C9.31C; C9.32C

## Progression questions

- How are carboxylic acids produced?
- What are the names, formulae and structures of the first four carboxylic acids?
- How does the functional group in all carboxylic acids influence their chemical properties?



A oxidising ethanol

If wine is left open to the air, it turns into vinegar due to a reaction between ethanol and oxygen. The product is **ethanoic acid**, which is responsible for the distinctive sharp taste of vinegar. This **oxidation** reaction, shown in diagram A, occurs naturally due to the presence of certain bacteria. Vinegar can be made from any dilute ethanol solution like wine, cider or beer.

- 1 What substance produces the sharp taste of vinegar?
- 2 Describe the loss and gain of atoms when an ethanol molecule is oxidised.

Ethanoic acid is a member of an homologous series called the **carboxylic acids**. The first four members of the series are shown in table C.

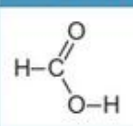
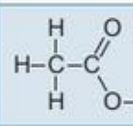
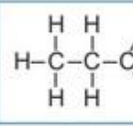
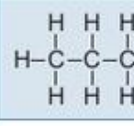
## Did you know?

Many chip shops do not use vinegar produced from fermented alcohol. Instead they use diluted ethanoic acid made from crude oil. It has exactly the same taste as the more expensive natural vinegar, but must be labelled as 'non-brewed condiment'.



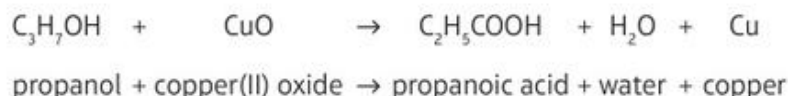
B

- 3 The fifth carboxylic acid is called pentanoic acid. Draw its structural formula and circle the functional group.

Name	Molecular formula	Structural formula
methanoic acid	HCOOH	
ethanoic acid	CH <sub>3</sub> COOH	
propanoic acid	C <sub>2</sub> H <sub>5</sub> COOH	
butanoic acid	C <sub>3</sub> H <sub>7</sub> COOH	

C the first four carboxylic acids

All carboxylic acids have the general formula  $C_nH_{2n+1}COOH$  and contain the  $-COOH$  functional group. They are often formed by the gentle oxidation of alcohols using **oxidising agents** such as hot copper oxide. For example:





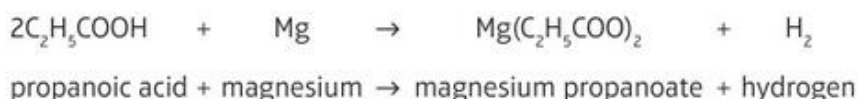
- 4** What carboxylic acid is formed by the oxidation of methanol?
- 5** Write word equations for the following oxidation reactions.
- a** methanol + copper(II) oxide
- b** ethanol + copper(II) oxide

The carboxylic acids have similar chemical properties. They all:

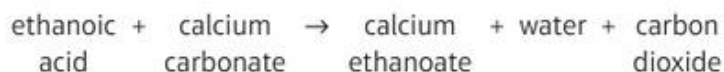
- form solutions with a pH less than 7 (if soluble)
- react with metals to form a salt and hydrogen
- react with bases to form a salt and water
- react with carbonates to form a salt, water and carbon dioxide.

The acidic properties are due to the presence of the  $\text{-COOH}$  group. In solution this group forms a hydrogen ion, while the rest of the molecule forms the negative ion that makes the salt. For example, propanoic acid forms propanoate ions, which make propanoate salts.

The equation for the reaction between magnesium and propanoic acid is:



Vinegar can be used to test for carbonate rocks like chalk and limestone (calcium carbonate). The ethanoic acid 'fizzes', producing carbon dioxide gas.



- 7** Write word equations for the following reactions.

- a** methanoic acid + lead(II) oxide
- b** butanoic acid + zinc carbonate
- 8** When ethanoic acid is added to sodium hydroxide solution the temperature increases.
- a** Name the products of this reaction.
- b** Suggest two possible names for this type of reaction.



**D** the first three carboxylic acids reacting with magnesium

- 6** Look at photo D.

- a** Write word equations for the three reactions.
- b** Suggest what the reaction between butanoic acid and magnesium would look like.

### Exam-style question

- a** Suggest names for the compounds with the formulae  $\text{C}_3\text{H}_7\text{OH}$  and  $\text{C}_2\text{H}_5\text{COOH}$ . (2 marks)
- b** Name the type of reaction that changes  $\text{C}_3\text{H}_7\text{OH}$  into  $\text{C}_2\text{H}_5\text{COOH}$ . (1 mark)

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** Construct a table showing the names, molecular formulae, functional groups and salts formed for the four simplest carboxylic acids.

### Extend

**E1** Compare and contrast alcohols with carboxylic acids.



# SC24a Addition polymerisation

Specification reference: C9.17C; C9.18C; C9.25C

## Progression questions

- What is a polymer?
- What monomers join together to form DNA, starch and proteins?
- How do ethene molecules join together to form poly(ethene)?





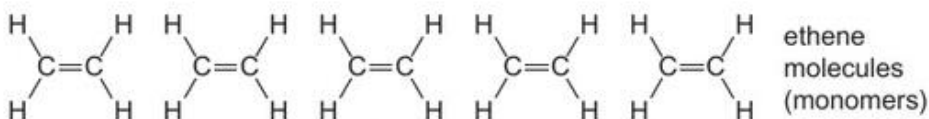
**A** Polytunnels, used for growing crops, are made from poly(ethene).

Plastics are made from **polymers**. A polymer is a large molecule made from lots of small molecules, called **monomers**, joined together. The process in which monomers join together is called **polymerisation**.

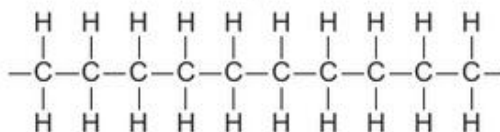
A polymer has a high average relative molecular mass, anything from tens of thousands to millions. The polymer molecules can be different lengths, so they cannot be given a specific relative molecular mass.

Poly(ethene), more commonly known as polythene, is a polymer formed when a large number of ethene monomers join together. Ethene molecules have a double covalent bond between the carbon atoms. One of the bonds in the double bond breaks open and another ethene molecule adds on. This process happens again and again, forming a long chain. The process in which monomers add together in this way is called **addition polymerisation**.

-  1 Describe what is meant by a polymer.
-  2 State the type of bonds between the carbon atoms in ethene.



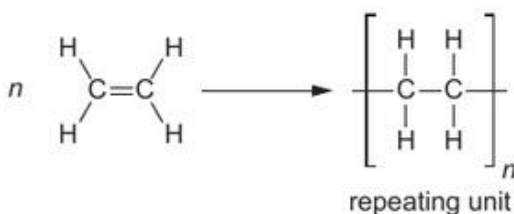
addition  
polymerisation



part of a  
poly(ethene)  
molecule  
(polymer)

**B** the formation of poly(ethene), shown using structural formulae

This can be written as an equation:



**D** In the equation for the formation of poly(ethene),  $n$  is a very large number. The repeating unit is shown in brackets with the subscript  $n$ .

## Did you know?





One of the first synthetic polymers was polyoxybenzylmethylenglycolanhydride. It was invented in 1907 by Leo Baekeland (1863–1944), and called Bakelite. Early telephones were made from it.



**C**

Equations for polymerisation show the structural formulae of the monomer and the **repeating unit**. The repeating unit shows how the monomer has changed and how it repeats throughout the polymer chain. The repeating unit does not have a double bond but has bond lines passing through the brackets to show that there is another repeating unit joined on each side.

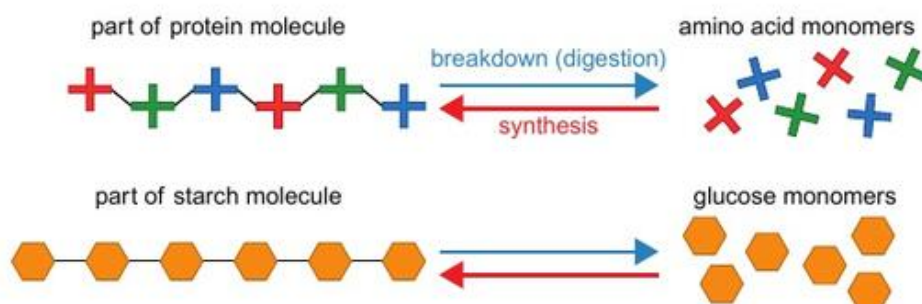
Poly(ethene) is a **synthetic polymer** as it is manufactured in a laboratory or factory.

-  **3** Describe how ethene molecules join together to form poly(ethene).
-  **4** State the type of polymerisation that takes place when ethene forms poly(ethene).
-  **5** Draw a part of a poly(ethene) molecule, showing two repeating units.
-  **6** Describe how the repeating unit in poly(ethene) differs from the monomer.




## Naturally occurring polymers

Some substances, such as DNA, starch and proteins, are **naturally occurring polymers**.

DNA is made from four different monomers called **nucleotides**. **Starch** is a polymer made from a **sugar** called glucose. **Proteins** are polymers made from **amino acids**.

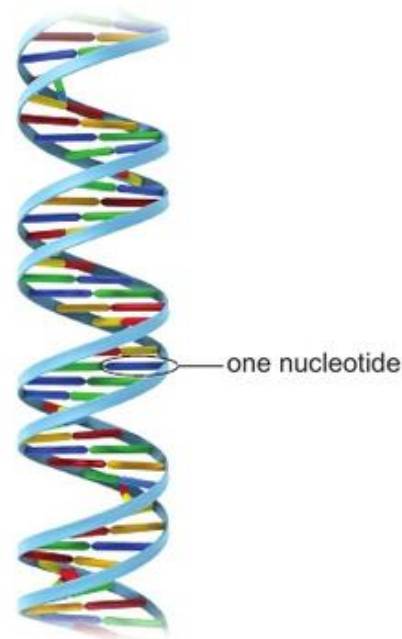


**F** Starch and proteins are polymers.

-  **7** Describe the difference between a synthetic polymer and a naturally occurring polymer, and give an example of each.
-  **8** State the name of the types of monomers that make up DNA.
-  **9** State the name of the naturally occurring polymer made from amino acids.

## Exam-style question

Write an equation to show the formation of poly(ethene) from ethene, showing the structures of the monomer and the polymer. (2 marks)



**E** part of a DNA molecule

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1** State the meanings of these terms: monomer, polymer, addition polymerisation.
- S2** State the names of the types of monomers that make up these polymers:  
**a** DNA   **b** starch   **c** protein.

## Extend

- E1** Explain, with the aid of diagrams, how two ethene molecules join together to form part of a poly(ethene) molecule.

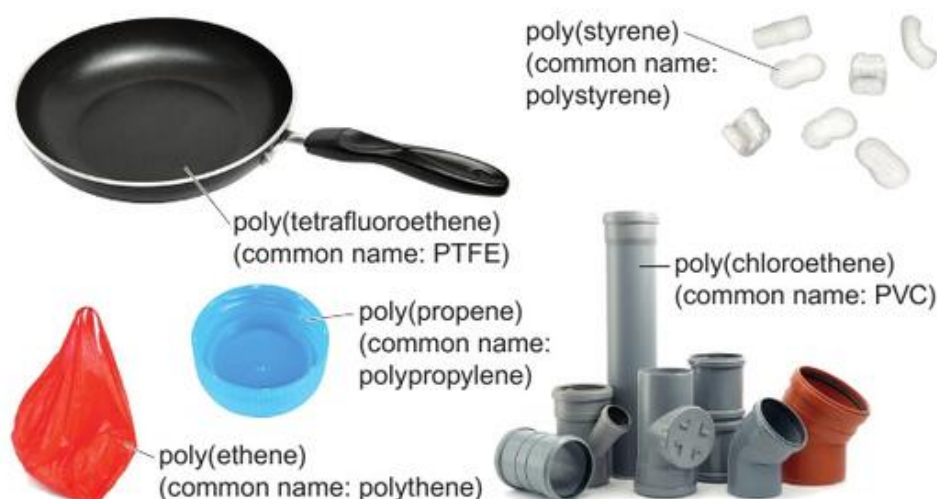


# SC24b Polymer properties and uses

Specification reference: C9.19C; C9.20C; C9.21C

## Progression questions

- How do chloroethene molecules join together to form poly(chloroethene)?
- How do you deduce the structure of a monomer from the structure of a polymer and vice versa?
- How are the uses of a polymer related to its properties?



The name of a polymer is in the form of 'poly' followed by the name of the monomer in brackets, where 'poly' means 'many'. Some polymers have common names as well, such as polythene for poly(ethene).

Poly(ethene) is formed when thousands of ethene molecules undergo addition polymerisation. Many other polymers can be made from alkene monomers because they all contain a C=C double bond. The general equation for any addition polymerisation is:

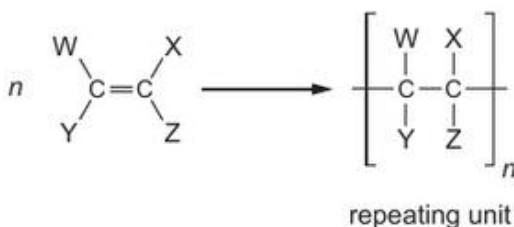
**A** There are many different polymers and their uses depend on their properties.



- 1** Give the name of the polymer formed from butene.



- 2** Give the name of the monomer used to form poly(methylpropene).



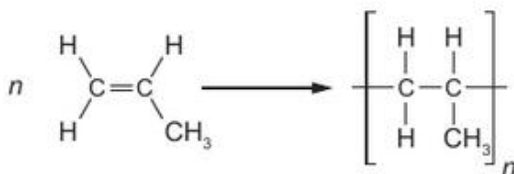
**B** In the general equation for addition polymerisation, W, X, Y and Z represent the atoms or groups of atoms attached to the C=C double bond in the monomer. Notice that these remain the same in the polymer.

Diagram C shows the equation for making poly(propene) from propene. There are only two carbon atoms in the carbon chain in the repeating unit. All the other atoms or groups of atoms that were attached to the carbons joined by the double bond remain in the same places.



- 3** Tetrafluoroethene has the formula  $\text{C}_2\text{F}_4$  and has a C=C double bond.

- a** Draw the structure of tetrafluoroethene, showing all the covalent bonds.



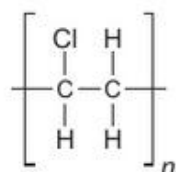
**C** equation for the formation of poly(propene)



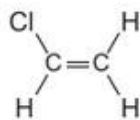
- b** Draw the repeating unit of poly(tetrafluoroethene).

You can work out the structure of the monomer from the formula for the polymer. Remove the brackets and the extension bond lines that go through them, and then draw a double bond between the two carbon atoms. Keep all the other atoms or groups of atoms the same.

For example, for poly(chloroethene):



repeating unit



monomer

**D** the structural formulae for the repeating unit and monomer of poly(chloroethene)

Polymers have different uses depending on their properties. Table F shows some common examples. New polymers are being developed all the time. For example, hydrogels are polymers that can absorb 500 times their own mass in water, making them useful for soft contact lenses and nappy linings.

Polymer	poly(ethene)	poly(propene)	poly(chloroethene)	poly(tetrafluoroethene)
Common name	polythene	polypropylene	polyvinyl chloride, PVC	PTFE, Teflon™
Properties	flexible, cheap, good insulator	flexible, does not shatter	tough, good insulator, can be made hard or flexible	tough, slippery
Uses	plastic bags, plastic bottles, cling film, polytunnels	buckets and bowls, crates, ropes, carpets	window frames, gutters, pipes, insulation for electrical wires	non-stick coatings for frying pans and kitchen utensils, burette taps, stain-proofing clothing and carpets

**F** some polymers and their uses

- 5** Select a polymer from table F that could be used to make each of the following items, and give a reason for your choice.



**a** a coating for a table that stops dust sticking to it



**b** a bucket and spade to use at the seaside

### Did you know?

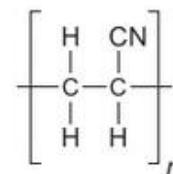
Old-fashioned waterproof coatings for clothing fabrics could keep water out but would not allow water vapour released as sweat to escape. This meant that the fabric still got wet on the inside. Modern polymer coatings for fabrics have now been developed that keep water out but also allow water vapour from sweat to escape so the wearer stays dry.

### Exam-style question

Draw a diagram to show part of a poly(propene) molecule formed from two propene molecules. The structure of the monomer is shown in diagram C. (2 marks)



- 4** A polymer, X, has the repeating unit:



**E** polymer X

Draw the structure of the monomer that formed polymer X.

### Checkpoint

How confidently can you answer the Progression questions?

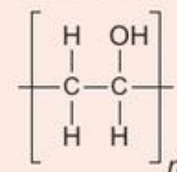
### Strengthen

**S1** Give reasons that explain why:

- a** PVC is used as a covering on electrical wires
- b** PTFE is used as a coating on frying pans.

### Extend

**E1** Draw the structure of the monomer used to form poly(ethanol). The repeating unit of the polymer is:



**G** poly(ethanol)



# SC24c Condensation polymerisation

Specification reference: **H** C9.22C

## Progression questions

- **H** What is meant by condensation polymerisation?
- **H** Which two functional groups react together to form a polyester?
- **H** How do you draw the structure of a polyester?

**H**



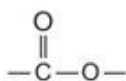
**A** Most plastic drinks bottles are made from PET.

Many plastic bottles are made from PET (poly(ethylene terephthalate)), which is a **polyester**. The bottles are strong, lightweight, shatter-resistant and can be recycled. They can be made into items such as plastic garden furniture or into fibres for use in items such as carpets and fleece jackets.

Polyesters are synthetic polymers made by **condensation polymerisation** in which the monomers join together and eliminate a small molecule, such as water.

- 6<sup>th</sup>** 1 State two properties of PET bottles.
- 6<sup>th</sup>** 2 State two items that can be made from recycled PET.
- 6<sup>th</sup>** 3 State what is meant by condensation polymerisation.

Organic compounds contain a **functional group**. A functional group is an atom or group of atoms that is responsible for the properties and reactions of the compound. The functional group in alcohols is  $\text{-OH}$  and in carboxylic acids it is  $\text{-COOH}$ . Esters are organic compounds that contain the functional group  $\text{-COO-}$ , which is drawn as:

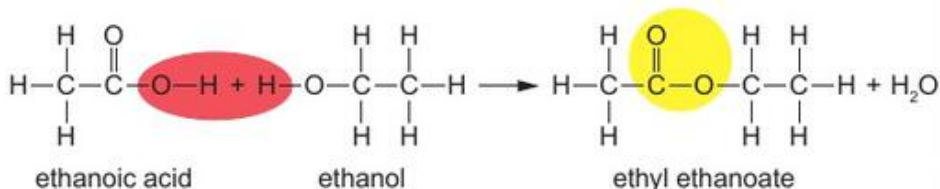


**B** the ester functional group

Esters are formed when a carboxylic acid reacts with an alcohol, in the presence of a catalyst. This is a condensation reaction because water is also produced.



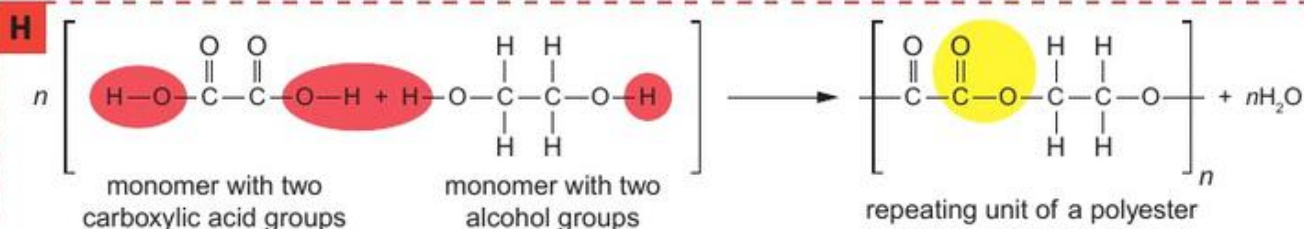
For example, ethanoic acid reacts with ethanol to form an ester, called ethyl ethanoate, and water.



- 6<sup>th</sup>** 4 Write the equation for the reaction between methanoic acid ( $\text{HCOOH}$ ) and methanol ( $\text{CH}_3\text{OH}$ ), showing the structures of the reactants and the product.

**C** This equation shows the formation of an ester from ethanoic acid and ethanol. The atoms in the red oval join together to form water. The yellow circle shows the ester link.

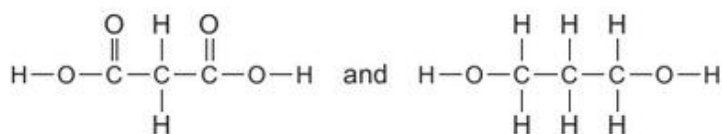
A polyester is a long-chain molecule that contains many **ester links** (in yellow in diagram C). The monomers that form a polyester contain alcohol and carboxylic acid functional groups. To form a long chain, each monomer must have two functional groups, one at each end of the molecule.



**D** This equation shows the formation of a polyester. The atoms shaded in red are eliminated from the monomers to form water. The group shaded in yellow is an ester link. Polyesters contain many ester links along the polymer chain.

Different polyesters can be formed by using monomers with different numbers of carbon atoms.

**5** A polyester is formed from the following monomers:



**E** polyester monomers



**a** Write the equation for the formation of this polyester.



**b** Highlight the ester link in the polyester.



**6** Explain how a molecule of water is formed each time an ester link is formed.

Polyesters are also used in the manufacture of synthetic fibres for weaving into fabrics for clothes and sheets.

Polyamides are another type of condensation polymer.



**F** Many items of clothing are made from polyesters or a mixture of polyesters and natural fibres such as cotton.

### Did you know?

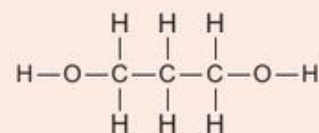
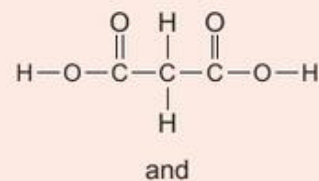
Nylon is a common synthetic polyamide formed by condensation polymerisation. Wallace Carothers (1896–1937) first produced it in 1935. Nylon was intended as a replacement for silk and was used to make parachutes and stockings during World War II. It is still used for parachutes but now has many other uses including cookware and car components.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1** Explain what is meant by condensation polymerisation.
- S2** Write the equation for the formation of a polyester from these monomers:



### Extend

- E1** Write the equation for the formation of the polyester formed from  $\text{HOOC}(\text{CH}_2)_2\text{COOH}$  and  $\text{HO}(\text{CH}_2)_2\text{OH}$ , showing all the bonds in the reactants and products.

### Exam-style question

State the functional groups that react together to form an ester link.

(1 mark)



# SC24d Problems with polymers

Specification reference: C9.23C; C9.24C

## Progression questions

- What problems are associated with making polymers?
- What problems are associated with the disposal of polymers?
- What are some advantages and disadvantages of recycling polymers?



**A** The extraction and transport of crude oil can have disastrous consequences for wildlife.

Crude oil is separated into fractions by **fractional distillation**. The fractions are further purified, **cracked** and refined to produce useful chemicals. Most of the monomers needed to make synthetic polymers are obtained from crude oil. Crude oil is a **finite resource** and is **non-renewable**, so chemists will need to find new sources of monomers in the future.

Materials such as wood and paper are **biodegradable**. This means that they rot because microbes can feed on them and break them down. Most synthetic polymers are useful for many purposes because they are not biodegradable and so they last for a long time. However, this also means that they do not rot when they are thrown away.



**B** These plastic items were found in the stomachs of dead albatrosses in Hawaii. The plastics are non-biodegradable and so they will not rot if they get into the sea. The albatrosses, such as the one in the photo, mistake the items for food.



**1** State the meaning of the term 'biodegradable'.

**2** Most synthetic polymers are non-biodegradable. State why this is:



**a** an advantage

**b** a disadvantage.

Over half of the rubbish in the UK goes to landfill sites. The non-biodegradable plastic materials in the rubbish will last for many years. Some waste is **incinerated** (burned) and the energy released can be used to generate electricity. However, all plastics produce carbon dioxide, a greenhouse gas, and some plastics produce toxic substances when they burn. Most of these toxic substances can be removed from the waste gases, but this forms toxic ash, which must be disposed of safely.

We can reduce the amount of waste that goes into incinerators or landfill sites by reducing the amounts of materials we use and also by reusing materials. For example, we can reuse plastic bags rather than just throwing them away after a single use. If an item cannot be reused any more, it may be possible to recycle it by processing it to make a new item. This will also help to conserve the supplies of crude oil.



**3** State three ways of disposing of plastics.



**4** Describe why incinerating synthetic polymers can cause problems.





## 5 Explain the difference between reusing and recycling.

It is difficult to recycle polymers because many steps are needed to obtain a new item. The main steps are outlined below.

- i Collect the waste – this may involve kerbside collection or people taking it to a collection point.
- ii Sort the waste into different types of polymer – this is often done by hand, which is time-consuming and expensive.
- iii Dispose of waste that cannot be recycled in a landfill site.
- iv Clean the polymers and grind them into chippings.
- v Purify the chippings.
- vi Melt the chippings then process them into a new product.

To make the sorting of polymers easier, codes and symbols are used on plastic items to show what they are made of. These are shown in table C. The majority of recycled items are those with recycling symbols 1 or 2.

symbol	polymer	uses
	poly(ethylene terephthalate)	some bottles, food trays, duvet fillings
	high-density poly(ethene)	some bottles, buckets
	poly(chloroethene)	soft toys, window frames
	low-density poly(ethene)	cling film, bags
	poly(propene)	crisp packets, carpet, rope
	poly(styrene)	egg boxes, foam packaging
	other polymers	

C polymer recycling symbols

## Did you know?

Chemists are developing new polymers that are biodegradable. For example, poly(lactic acid) is made from cornstarch.



D a biodegradable bag will rot

## 6 Identify the steps in recycling polymers that require energy. State what the energy is used for.

## Exam-style question

Polymers persist for a long time in landfill sites. State *one* way in which this problem can be overcome. (1 mark)

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

- S1 Describe some problems associated with:
  - a the starting materials for polymers
  - b the disposal of polymers.
- S2 Describe the advantages and disadvantages of disposing of plastics by:
  - a putting them in landfill sites
  - b burning them.

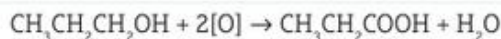
## Extend

- E1 Evaluate the advantages and disadvantages of recycling polymers.



## Alcohols and carboxylic acids

When heated together, acidified potassium dichromate solution oxidises propanol to form propanoic acid. The reaction can be modelled using this balanced equation (where 2[O] represents two oxygen atoms added in the reaction):



Propanol and ethanol are members of the alcohol homologous series. Propanoic acid and ethanoic acid are members of the carboxylic acid homologous series.

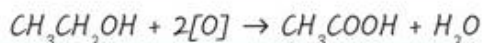
Describe how ethanoic acid can be produced from sugar. The first step is to obtain a concentrated solution of ethanol. You may use equations if you wish. (6 marks)

### Student answer

Dissolve the sugar in warm water, then add yeast. Yeast enzymes catalyse fermentation. The reaction mixture must be kept at 25–35 °C so that the enzymes work at their best. An air lock is needed so that carbon dioxide can escape, but air cannot get in [1].

Fermentation takes several days. When it is complete, remove the yeast by filtration [2]. Separate ethanol from the filtrate using fractional distillation. This works because ethanol has a lower boiling point than water, so it is collected first [3].

Use the concentrated ethanol solution to make ethanoic acid. Heat the ethanol with acidified potassium dichromate solution:



This works because ethanol and propanol are both alcohols. They have the same functional group and similar chemical properties [4].

[1] The answer explains the temperature needed. It could also mention that fermentation is an anaerobic process, which is why air must be kept out.

[2] The reason for filtering the reaction mixture is given.

[3] The answer explains why a concentrated solution of ethanol can be produced by fractional distillation.

[4] The balanced equation given in the question is applied to predict the reaction of ethanol.

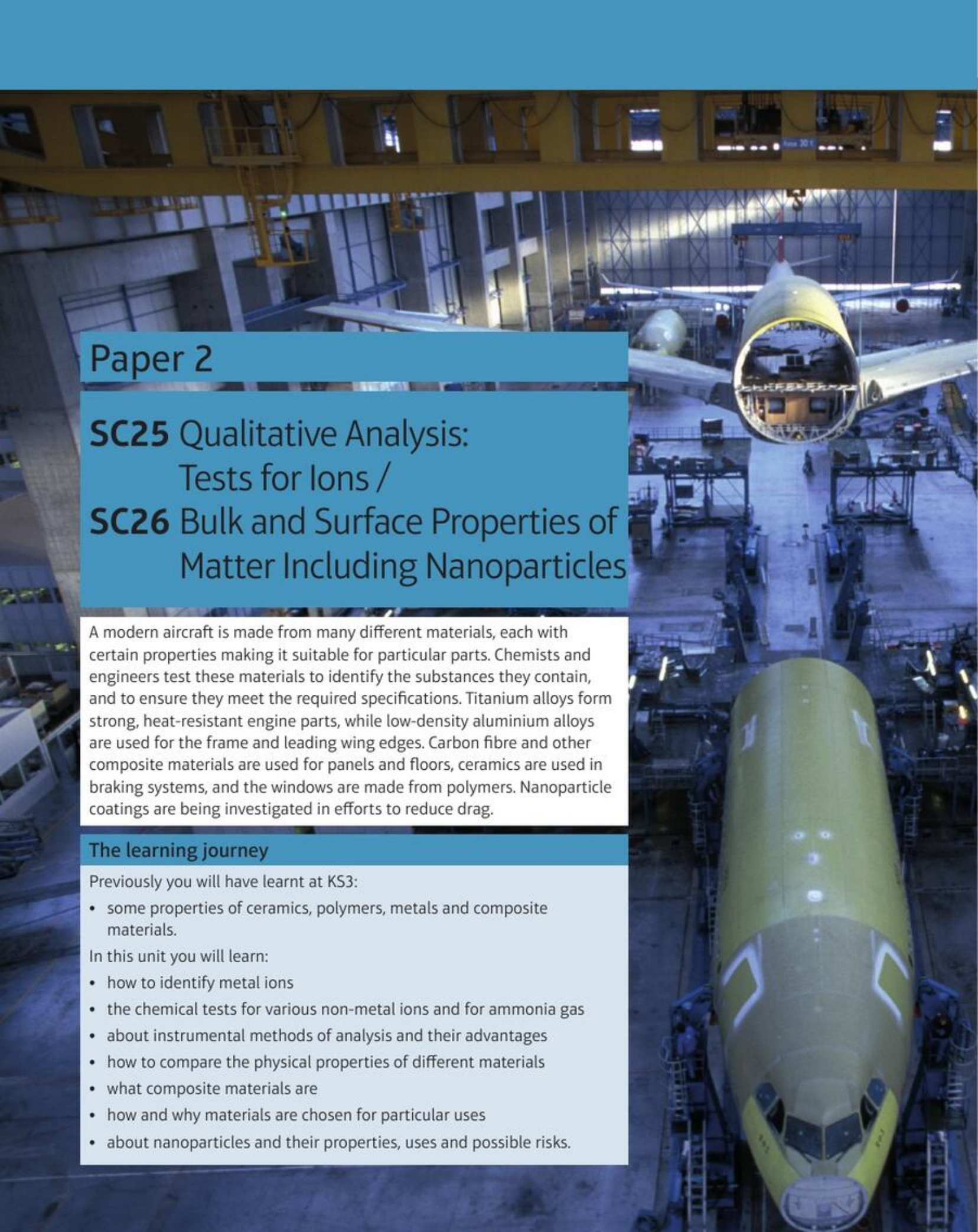
### Verdict

This is a strong answer. It shows good knowledge and understanding of the production of ethanol by fermentation, how to produce a concentrated solution of ethanol and the similarity in chemical properties between members of an homologous series. The answer is presented in a logical order with good use of appropriate terminology.

#### Exam tip

Make sure you are prepared to apply information and so can, for example, predict the reactions of other members of a given homologous series.



A large commercial airplane is shown in a hangar with its nose open, revealing the cockpit area. The hangar is filled with various equipment and structural elements, and the lighting is bright, highlighting the aircraft's white exterior and the interior of the nose.

## Paper 2

### **SC25** Qualitative Analysis: Tests for Ions / **SC26** Bulk and Surface Properties of Matter Including Nanoparticles

A modern aircraft is made from many different materials, each with certain properties making it suitable for particular parts. Chemists and engineers test these materials to identify the substances they contain, and to ensure they meet the required specifications. Titanium alloys form strong, heat-resistant engine parts, while low-density aluminium alloys are used for the frame and leading wing edges. Carbon fibre and other composite materials are used for panels and floors, ceramics are used in braking systems, and the windows are made from polymers. Nanoparticle coatings are being investigated in efforts to reduce drag.

#### The learning journey

Previously you will have learnt at KS3:

- some properties of ceramics, polymers, metals and composite materials.

In this unit you will learn:

- how to identify metal ions
- the chemical tests for various non-metal ions and for ammonia gas
- about instrumental methods of analysis and their advantages
- how to compare the physical properties of different materials
- what composite materials are
- how and why materials are chosen for particular uses
- about nanoparticles and their properties, uses and possible risks.



# SC25a Flame tests and photometry

Specification reference: C9.2C; C9.7C; C9.8C; C9.9C

## Progression questions

- How are metal ions identified using flame tests?
- Why might chemists analyse substances using machines instead of chemical tests?
- How is the information from flame photometers used?



**A** The yellow light from these street lamps is emitted when an electric current passes through a vapour of sodium ions.

Flame tests are used to identify metal ions in substances. To carry out a flame test:

- i Light a Bunsen burner and open the air hole to give a hot blue flame.
- ii Pick up a small sample of the test substance using a wire loop.
- iii Hold the sample in the edge of the flame and observe the flame colour.

The wire loop is cleaned in hydrochloric acid before testing each sample. Platinum wire has a high melting point and it is unreactive. It also gives no colour to the flame. However, nichrome alloy is often used instead. Although it produces its own faint orange colour, it is much cheaper.



**B** Sodium ions produce a yellow flame test colour.



- 1** Wooden splints may be used instead of a wire loop. Suggest two reasons that explain why the splints are soaked in distilled water before use.

Different metal **cations** produce different flame test colours (as shown in table C). Flame tests work with solid samples and with solutions, but flame colours from solids are easier to see.

Metal ion	Symbol	Flame test colour
lithium	$\text{Li}^+$	red
sodium	$\text{Na}^+$	yellow
potassium	$\text{K}^+$	lilac
calcium	$\text{Ca}^{2+}$	orange-red
copper	$\text{Cu}^{2+}$	blue-green

**C** some flame test colours



- 2** Explain why potassium chloride and potassium iodide both produce lilac flame test colours.

- 3** Predict the flame test colours produced by the following compounds.

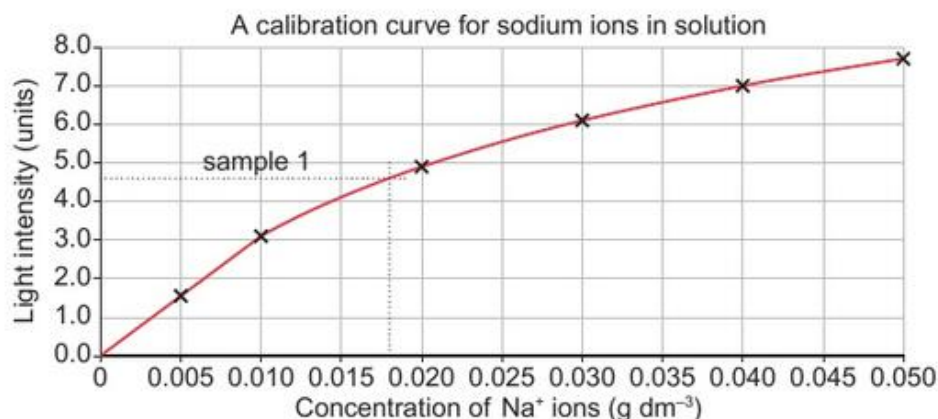


- a**  $\text{CaCl}_2$     **b**  $\text{Na}_2\text{SO}_4$     **c**  $\text{LiNO}_3$     **d**  $\text{Cu}_3(\text{PO}_4)_2$

## Flame photometry

Machines can also be used to analyse substances. Compared to simple laboratory tests such as flame tests, using scientific instruments may improve:

- sensitivity (they can detect much smaller amounts)
- accuracy (they give values closer to the true values)
- speed.



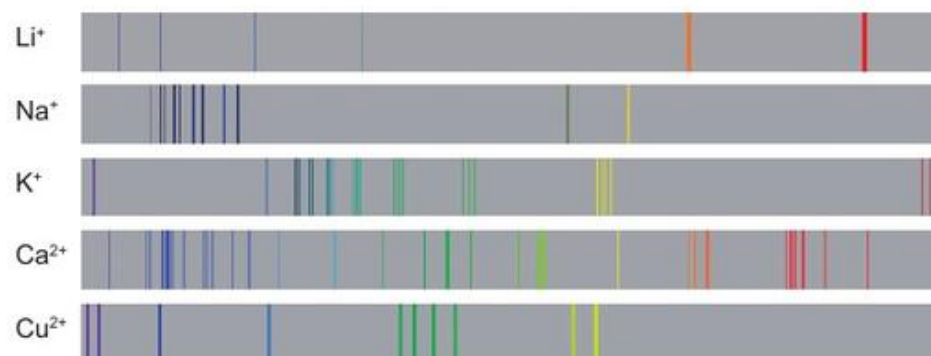
D a calibration curve from a flame photometer

The **flame photometer** measures the light intensity of the flame colours produced by metal ions. Its data is used to determine the concentration of a metal ion in a dilute solution. Graph D shows a **calibration curve** produced using different **standard solutions**, each containing known concentrations of the metal ion dissolved in distilled water.

- 5 In graph D, sample 1 gives a light intensity reading of 4.6 units, so its sodium ion concentration is  $0.018 \text{ g dm}^{-3}$ . A second sample gives a light intensity reading of 6.1 units. Determine its sodium ion concentration.

- 4 Describe the relationship between light intensity and concentration shown in graph D.

The colour of light that we see in a flame test is usually a mixture of different colours. A flame photometer can separate out these colours to produce a **spectrum** of the light emitted by each metal ion. Different metal ions produce different **emission spectra** (shown in diagram E). A metal ion in an unknown solution can be identified by matching its spectrum to the spectrum from a known metal ion.



E emission spectra for some metal ions

- 6 Compare and contrast the emission spectra shown in diagram E.

### Did you know?

Robert Bunsen and Gustav Kirchhoff analysed the spectrum of light given off by heated mineral salts. Using this method, they discovered caesium in 1860 and rubidium in 1861.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

- S1 How are metal ions in solids identified using flame tests?
- S2 How are the data from instrumental methods of analysis used?

### Extend

- E1 Evaluate the use of flame tests compared to the use of a flame photometer.

### Exam-style question

Explain why a flame test could *not* be used to identify the metal ions in a mixture of lithium chloride and calcium chloride. (2 marks)



# SC25b Tests for positive ions

Specification reference: C9.1C; C9.3C; C9.4C; C9.7C

## Progression questions

- Why must the test for an ion only detect that ion?
- How are metal ions identified using sodium hydroxide solution?
- How are ammonium ions and ammonia detected?



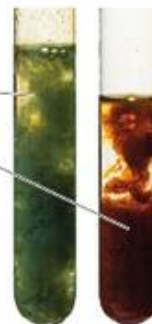
**A** This stream is discoloured by metal ions in waste water draining from a mine.

Metal compounds dissolve in groundwater as it flows through mines. When it drains from the mines, the water contaminates rivers. One way to treat the water involves sodium hydroxide solution. This reacts with metal ions to produce **precipitates**, which settle out and can be removed.

## Testing for metal ions

**Precipitation reactions** involving sodium hydroxide solution form the basis of a test to identify dissolved metal ions. A few drops at a time of dilute sodium hydroxide solution are added to the test solution. Different metal ions produce different coloured hydroxide precipitates, as shown in table B.

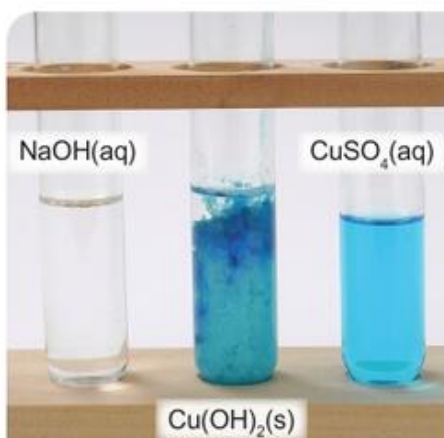
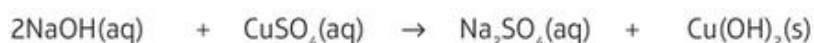
Metal ion	Symbol	Precipitate colour
iron(II)	$\text{Fe}^{2+}$	green
iron(III)	$\text{Fe}^{3+}$	brown
copper	$\text{Cu}^{2+}$	blue
calcium	$\text{Ca}^{2+}$	white
aluminium	$\text{Al}^{3+}$	white



**B** metal hydroxide precipitate colours for different metal ions

For example, sodium hydroxide solution reacts with copper sulfate solution to form a blue precipitate:

sodium hydroxide + copper sulfate  $\rightarrow$  sodium sulfate + copper hydroxide



**C** A blue precipitate of copper hydroxide forms when sodium hydroxide solution reacts with copper sulfate solution.



- 1 A few drops of sodium hydroxide solution are added to a colourless solution. A faint green precipitate forms. Identify the metal ions present in the colourless solution.

- 2 For the reaction between sodium hydroxide solution and iron(II) chloride solution:



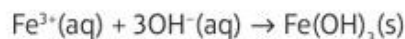
**a** write the word equation



**b** write the balanced equation, including state symbols.

**H**

These precipitation reactions can also be modelled using balanced ionic equations. For example, aqueous iron(III) ions react with aqueous hydroxide ions to form solid iron(III) hydroxide:



- 3** Write the balanced ionic equation, including state symbols, for the formation of calcium hydroxide,  $\text{Ca}(\text{OH})_2$ .

It is important that the test for an ion should only detect that ion. For example, calcium ions and aluminium ions both produce white hydroxide precipitates. You cannot distinguish between these without a further test. Aluminium hydroxide disappears to form a colourless solution when excess sodium hydroxide solution is added, but calcium hydroxide does not.



- 4** Explain how you could use dilute sodium hydroxide solution to distinguish between calcium sulfate solution and aluminium sulfate solution.

### Testing for ammonium ions

Dilute sodium hydroxide solution is also used to detect ammonium ions,  $\text{NH}_4^+$ , in a substance. Ammonia gas,  $\text{NH}_3$ , is produced when the mixture containing ammonium ions is warmed. Ammonia has a characteristic sharp smell, but a **confirmatory test** is used to identify it. Ammonia changes the colour of damp red litmus paper to blue.



- 5** Explain the meaning of the colour change seen when damp red litmus paper is used to test for ammonia.



- 6** **H** Write a balanced ionic equation to show the reaction between aqueous ammonium ions and aqueous hydroxide ions, producing ammonia gas and one other product. Include state symbols.

### Did you know?

Aluminium sulfate is used to treat fresh water to make it potable (safe for drinking). Aluminium ions react with other ions in the water to produce an aluminium hydroxide precipitate. This catches tiny particles, such as rotted plant material and clay, forming a solid that sinks to the bottom of the treatment tank.



**D** In another test for ammonia, a glass rod dipped in concentrated hydrochloric acid is held near the mouth of the test tube. A white smoke of ammonium chloride forms.

### Checkpoint

How confidently can you answer the Progression questions?

### Strengthen

**S1** How is sodium hydroxide solution used to identify positively charged ions?

### Extend

**E1** To what extent can sodium hydroxide solution be used to identify all positively charged ions?

### Exam-style question

Ammonium iron(III) sulfate is a soluble solid used in waste water treatment. Describe how you could show that it contains ammonium ions and iron(III) ions. (4 marks)



# SC25c Tests for negative ions

Specification reference: C9.5C; C9.7C

## Progression questions

- How are carbonate ions and carbon dioxide detected?
- How are sulfate ions detected?
- How are halide ions identified?






**A** This geologist is testing tufa, a type of limestone rock, with dilute acid.

Geologists carry out simple 'acid tests' on rocks. If a rock contains carbonate minerals, it produces bubbles when dilute hydrochloric acid is added to it.

## Testing for carbonate ions

Just as there are simple chemical tests to detect and identify positively charged ions (cations), there are tests for negatively charged ions (**anions**). Carbonate ions,  $\text{CO}_3^{2-}$ , are easily detected in solids and in solutions. Add dilute hydrochloric acid to the test substance and look for bubbling caused by the production of carbon dioxide. Since the bubbles could be due to a different gas, such as hydrogen, you should carry out a confirmatory test. If the bubbles do contain carbon dioxide, they turn limewater milky.

-  **1** Describe how to detect carbonate ions in sodium carbonate solution.
-  **2**  Write a balanced ionic equation to show the reaction between aqueous hydrogen ions (produced by acids in solution) and aqueous carbonate ions to form carbon dioxide gas and water. Include state symbols.

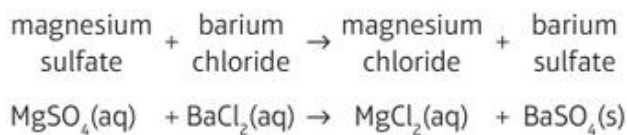


**B** using limewater to confirm the results of a positive test for carbonate ions

## Testing for sulfate ions

To detect sulfate ions in a solution, add a few drops of dilute hydrochloric acid. This acidifies the solution and removes carbonate ions that might also give a precipitate in the test. Then add a few drops of barium chloride solution. A white precipitate of barium sulfate forms if the sample contains sulfate ions,  $\text{SO}_4^{2-}$ .

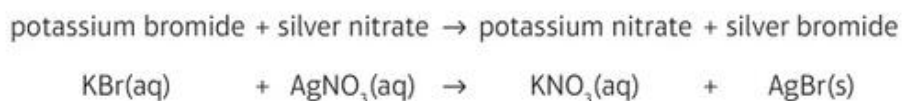
For example:



- 3** Suggest a reason that explains why dilute sulfuric acid,  $\text{H}_2\text{SO}_4$ , is *not* used to acidify the sample when testing for sulfate ions.
- 4** Write an ionic equation, including state symbols, for the reaction between barium ions and sulfate ions.

## Testing for halide ions

The halogens are the non-metal elements in group 7 of the periodic table. They form compounds that contain **halide ions**, such as  $\text{F}^-$ ,  $\text{Cl}^-$ ,  $\text{Br}^-$  and  $\text{I}^-$ . Silver fluoride is soluble in water but the other silver **halides** are insoluble. This is the basis of the laboratory test for chloride, bromide and iodide ions. To detect these ions in a solution, add a few drops of dilute nitric acid. This acidifies the solution and removes carbonate ions that might also give a precipitate in the test. Then add a few drops of silver nitrate solution. Different coloured silver halide precipitates form, depending on the halide ion present (see table D and photo E). For example:



Halide ion	Symbol	Precipitate colour
chloride	$\text{Cl}^-$	white
bromide	$\text{Br}^-$	cream
iodide	$\text{I}^-$	yellow

**D** silver halide precipitate colours

- 5** Explain why fluoride ions cannot be detected in the same way as chloride, bromide and iodide ions.
- 6** Substance X gives an orange-red flame test result and produces a yellow precipitate when silver nitrate solution is added. Name substance X.
- 7** Write an ionic equation for the reaction between silver ions and bromide ions. Include state symbols.

## Exam-style question

Silver carbonate is an insoluble yellow solid. Suggest an explanation for why dilute acid must first be added when testing solutions for halide ions using silver nitrate solution. (3 marks)



**C** A white precipitate of barium sulfate forms in a positive test for sulfate ions.



**E** silver halide precipitates

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** How are negatively charged ions identified in chemical tests?

## Extend

**E1** Explain why different dilute acids are used in the tests for negatively charged ions.



# SC25c Core practical – Identifying ions

Specification reference: C9.6C

## Aim

Identify the ions in some unknown salts, using the tests for the specified cations and anions.

Analytical chemists use different techniques to determine which substances are present or absent.

## Your task

You will use laboratory tests to identify the cations and anions in some unknown salts. You can then work out the name of each salt.



**A** Analytical chemists work in many areas, including chemical and forensic analysis, quality control, medical drug development and toxicology.

## Methods

For all tests, wear eye protection and avoid skin contact with all the chemical substances.

### Flame tests for metal cations

- A** Light a Bunsen burner and open the air hole to give a hot blue flame.
- B** Pick up a small sample of a solid salt using a clean wire loop or a damp wooden splint.
- C** Hold the sample in the edge of the flame.
- D** Observe and record the flame colour.

### Hydroxide precipitate tests for metal cations

- E** Dissolve a little solid salt in a test tube using distilled water.
- F** Add a few drops of dilute sodium hydroxide solution, one drop at a time.
- G** Record the colour of any precipitate formed.
- H** If a white precipitate forms, add excess dilute sodium hydroxide solution to see if it will disappear to leave a clear solution.

### Testing for ammonium ions

- I** Dissolve a little solid salt in a test tube using distilled water.
- J** Add dilute sodium hydroxide solution and then warm gently.
- K** Remove from the flame. Hold a piece of damp red litmus paper near the mouth of the test tube. Record what happens to its colour.

### Testing for carbonate ions

- L** Put a little solid salt in a test tube and add a few drops of dilute acid.
- M** Record whether any effervescence (bubbling) occurs.
- N** Use limewater to check that any bubbles contain carbon dioxide.

### Testing for sulfate ions

- O** Dissolve a little solid salt in a test tube using distilled water.
- P** Add a few drops of dilute hydrochloric acid, then a few drops of barium chloride solution. Record whether a white precipitate forms.

### Testing for halide ions

- Q** Dissolve a little solid salt in a test tube using distilled water.
- R** Add a few drops of dilute nitric acid, then a few drops of silver nitrate solution. Record the colour of any precipitate formed.

## Exam-style questions

- Copy diagram B. On your diagram, draw one straight line from each metal cation to its flame test colour. (5 marks)
- Some metal cations react with sodium hydroxide solution to form metal hydroxide precipitates.
  - Copy and complete table C. (3 marks)
  - Describe a test to distinguish between aluminium hydroxide and calcium hydroxide. (2 marks)
- Halide ions can be identified using silver nitrate solution. Copy table D. Place a tick (✓) in one box of each row to show the colour of the silver halide precipitate formed. (3 marks)
- A teacher demonstrates how to carry out flame tests using a platinum wire loop.
  - Give two reasons that explain why platinum is a suitable metal to use. (2 marks)
  - Give a reason that explains why a luminous Bunsen burner flame is *not* suitable for flame tests. (1 mark)
  - The teacher uses hydrochloric acid to clean the wire loop between each test. Explain why the teacher cleans the wire loop. (2 marks)
- Describe the laboratory test for carbon dioxide. (2 marks)
- A student tests a solution of an unknown salt. He adds a few drops of dilute nitric acid followed by a few drops of silver nitrate solution.
  - Give a reason that explains why he adds dilute nitric acid. (1 mark)
  - Explain why he should *not* use dilute hydrochloric acid instead of dilute nitric acid. (2 marks)
- Barium chloride is toxic.
  - Suggest a reason that explains why *dilute* barium chloride solution, rather than concentrated barium chloride solution, is used in tests to identify sulfate ions. (1 mark)
  - State and explain a suitable precaution to reduce the risk of harm when carrying out tests for sulfate ions. (2 marks)
- A solution of substance X gives a yellow flame test result. It produces a white precipitate when a few drops of dilute nitric acid are added followed by silver nitrate solution. Name substance X. (2 marks)
- A solution of substance Y produces a green precipitate when sodium hydroxide solution is added. It produces a white precipitate when dilute hydrochloric acid is added followed by barium chloride solution.
  - Name substance Y. (2 marks)
  - On standing, the green precipitate gradually turns brown. Suggest a reason that explains this observation. (1 mark)

Li <sup>+</sup>	•
Na <sup>+</sup>	•
K <sup>+</sup>	•
Ca <sup>2+</sup>	•
Cu <sup>2+</sup>	•

•	orange-red
•	red
•	blue-green
•	yellow
•	lilac

B

Metal cation	Colour of metal hydroxide
aluminium, Al <sup>3+</sup>	white
calcium, Ca <sup>2+</sup>	white
copper, Cu <sup>2+</sup>	
iron(II), Fe <sup>2+</sup>	
iron(III), Fe <sup>3+</sup>	

C

Halide ion	Colour of silver halide		
	white	yellow	cream
chloride, Cl <sup>-</sup>			
bromide, Br <sup>-</sup>			
iodide, I <sup>-</sup>			

D



# SC26a Choosing materials

Specification reference: C9.38C; C9.39C

## Progression questions

- What are ceramics?
- What are ceramics, polymers and metals like?
- How are materials chosen for a given use?



**A** Bathrooms contain many different materials, including glass and clay ceramics, polymers and metals.



**B** These new ceramic toilet fittings have just come out of a kiln, a type of oven.



**C** a modern window glass factory

## Glass and clay ceramics

**Ceramics** are a range of durable compounds that change very little when heated. They are chemically unreactive, hard and stiff but brittle. They are also poor electrical and thermal conductors, and have high melting points. Ceramic materials consist of giant structures with many strong bonds (covalent or ionic), giving them their typical properties.



- 1** State the typical physical properties of ceramic materials.

Brick, porcelain and china are **clay ceramics**. They are made from clay moulded into the desired shape. When the clay is heated to a very high temperature, tiny crystals form and join together. Bricks are usually decorated by adding a coloured substance to the clay before heating. A pattern may also be moulded into surfaces that will be visible in a finished wall. Porcelain and china are dipped in a 'glaze' and heated strongly again. The glaze forms the hard, waterproof, smooth surface you see on tiles, washbasins and toilet bowls.



- 2** Describe two properties of glazed clay ceramics that make them suitable for toilet bowls.

**Glass** is made by melting sand, then allowing it to cool and solidify. Glass and clay ceramics have similar properties because they both have giant structures. However, the atoms in glass are not arranged in a regular way to form crystals, so glass is **transparent** rather than **opaque**.

Modern window glass is made by the float process. Molten glass is poured onto a bath of molten tin, where it spreads out on the surface. The flat layer of glass is drawn away and cooled in a continuous process.



- 3** Describe two properties of glass that make it suitable for shower screens.



**Did you know?**

Medieval glass windowpanes are often thicker at the bottom, but not because glass flows very slowly. Instead, it is a result of the way the glass was made. A disc of molten glass was spun to make it spread out thinly, then cut into shape. It was installed thicker end down to make it more stable in the window frame.

**Polymers**

**Polymers** are substances with high average relative formula masses. They are made from **monomers** – smaller molecules that join together to form repeating units. For example, poly(ethene) is made from ethene, and poly(chloroethene), or PVC, is made from chloroethene (see SC24 *Polymers*).

Polymers can be moulded into complex shapes. The properties of a polymer depend on its structure and chemical composition, but polymers are usually strong and chemically unreactive. They are also poor electrical and thermal conductors. Rigid PVC is useful for underground pipes and window frames. PVC can be made softer by including substances called **plasticisers** in its manufacture. Flexible PVC is useful for indoor water pipes and waterproof flooring.

**Metals**

Most metals are strong, hard, shiny solids with high melting points (see SC7 *Types of Substance*). They are good electrical and thermal conductors. They are also **malleable**. Different metal elements can be mixed together to form **alloys**. The alloys are often stronger than the individual metals (see SC13 *Transition Metals, Alloys and Corrosion*). Metals can also be electroplated with an unreactive metal to improve their appearance or their resistance to corrosion.

- 5** Explain why bathroom taps are usually made from a metal electroplated with chromium.
- 6** Explain why bathroom mirrors are made from a sheet of glass coated on the back with a layer of aluminium.

**Exam-style question**

The table shows some information about copper and PVC, a flexible polymer.

Material	Relative electrical conductivity	Tensile strength (MPa) (resistance to stretching)
copper	1 000 000	220
PVC	1	22

Explain, using information from the table, why electrical cables are made from copper surrounded by a layer of PVC. (3 marks)



**D** PVC and metal plumbing fittings

- 4** Describe two properties of PVC that make it suitable for making toilet seats.

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

**S1** How are materials chosen for use in a bathroom?

**Extend**

**E1** Evaluate the use of different materials for a bench at a bus stop.



# SC26b Composite materials

Specification reference: C9.38C; C9.39C

## Progression questions

- What are composite materials?
- What are composite materials like?
- How are materials, including composite materials, chosen for a given use?

**6th** 1 Magnalium is an alloy made from a mixture of magnesium and aluminium. Explain why magnalium is *not* a composite material.

**7th** 2 Proposals for a pykrete aircraft carrier were put forward in the 1940s. Suggest an explanation for why this was impractical.

A **composite material** is a mixture of two or more materials, combined to produce a material with improved properties. The individual materials often have contrasting properties. The individual materials are also usually visible in the composite material, and can often be separated out by physical separation methods. Pykrete consists of ice and about 14% wood pulp. Tiny pieces of wood can be seen in pykrete, and they separate out when the ice melts.



A The base of this ice dome is made from pykrete. Pykrete is almost half as strong as concrete but much more lightweight.

## Reinforcement and matrix

Concrete is made by mixing cement, sand, aggregate (small stones and gravel) and water together. As the concrete sets hard, chemical reactions happen that bond the solid components together. The sand and aggregate form the **reinforcement** of the concrete. The reinforcement is bonded together by cement, which forms the **matrix**.

Many composite materials are like this. For example, in fibreglass, thin glass fibres form the reinforcement and a polymer resin forms the matrix. Glass fibres are brittle but have a low density and a high **tensile strength** (they resist being stretched). The polymer resin is hardwearing but not strong. The combination of these properties produces a lightweight, strong and tough composite material.



B These bicycle forks are made from a carbon-fibre-reinforced polymer. A liquid polymer resin is applied to a woven fabric made from carbon fibres. The resin sets hard and holds the fabric in shape.

**6th** 3 Name the reinforcement and matrix in a carbon-fibre-reinforced polymer.



Concrete can be reinforced with steel. Table C shows some typical properties of concrete and steel. **Compressive strength** is a measure of how well a material resists being squashed.

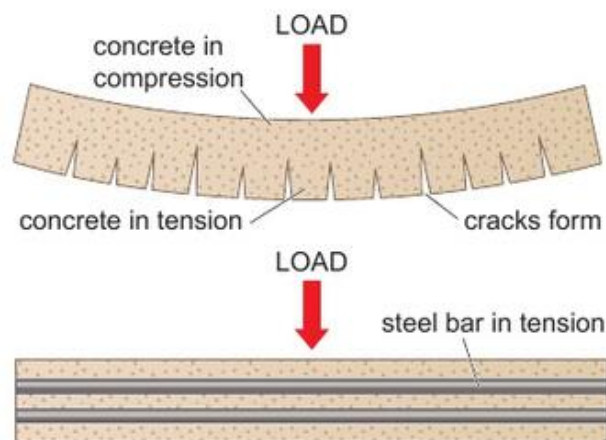
Material	Tensile strength (MPa)	Compressive strength (MPa)	Density (kg/m <sup>3</sup> )
concrete	4	40	2300
steel	500	300	7700

**C** some properties of concrete and steel

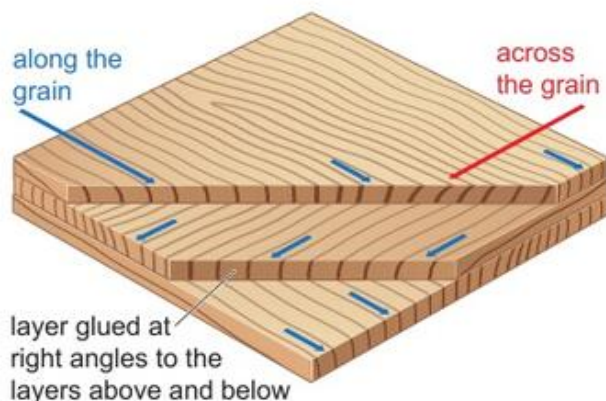
Concrete is strong in compression but weak in tension. It is suitable for road surfaces and for foundations, where it will mainly be squashed. However, concrete beams tend to crack along their lower surface, which is stretched when a beam carries a load (see diagram D). Steel is denser than concrete but has a much higher tensile strength. Steel-reinforced concrete consists of a framework of steel bars with concrete poured in and around it. The composite material formed is strong in tension *and* in compression, and is less dense and cheaper than steel alone.

## Laminates

Wood is a natural composite material consisting of cellulose fibres in a matrix of a polymer called lignin. It is stronger along its grain than it is across its grain. Plywood typically consists of odd numbers of thin sheets of wood, each glued at right angles to the sheet below (see diagram E).






**D** Steel-reinforced concrete resists cracking much better than concrete alone.



**E** In plywood, the glue that sticks the layers together forms part of the composite material.

Material	Tensile strength (MPa)
wood (along the grain)	80
wood (across the grain)	4
plywood (various directions)	25–40

**F** example wood and plywood strengths

-  **4** Explain, using information from table C, which material is stronger when stretched than when it is squashed.
-  **5** Suggest reasons that explain why steel should not be exposed at the surface of steel-reinforced concrete.
-  **6** Compare and contrast plywood with wood. Use information from table F in your answer.

## Exam-style question

Explain the meaning of the term 'composite material'.

(2 marks)

## Checkpoint

How confidently can you answer the Progression questions?

## Strengthen

**S1** What are the advantages of building a road bridge from steel-reinforced concrete, rather than from steel or concrete alone?

## Extend

**E1** Evaluate the use of papier mâché, expanded polystyrene or clear polymer made from recycled drinks bottles, to make egg boxes.

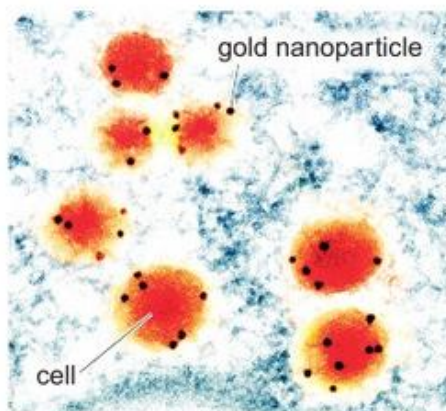


# SC26c Nanoparticles

Specification reference: C9.35C; C9.36C; C9.37C

## Progression questions

- Why do nanoparticulate materials have different properties from bulk materials?
- What are some of the uses of nanoparticles?
- What are some of the possible risks from nanoparticles?



**A** 'Immunogold labelling' allows biologists to investigate the different types of cells in a tissue. Gold nanoparticles are joined to antibody molecules, which then attach to the surfaces of certain cells.

Lumps and powders are examples of **bulk** materials, and consist of huge numbers of atoms. **Nanoparticles** typically consist of just a few hundred atoms, and are 1–100 nm in size. Nanoparticles are larger than atoms (e.g. an oxygen atom is 0.1 nm in diameter) and simple molecules. Nanoparticles are smaller than cells (e.g. a typical bacterium is 1000 nm long).

## Worked example W1

A gold nanoparticle is 32 nm in diameter.

- a** Calculate its diameter in metres, m.

$$1 \text{ nm} = 1 \times 10^{-9} \text{ m}$$

$$32 \text{ nm} = 32 \times 10^{-9} \text{ m} = 3.2 \times 10^{-8} \text{ m}$$

- b** The diameter of a gold atom is 0.28 nm. Estimate how many times larger the gold nanoparticle is compared to a gold atom.

Rounding each number to 1 significant figure gives 30 nm and 0.3 nm.

$$\text{Number of times larger} \approx \frac{30}{0.3} = 100$$



**B** Michael Faraday (1791–1867) prepared the first pure sample of nanoparticulate gold in 1857. The pink colour in Faraday's nanoparticulate gold is caused by the way nanoparticles scatter light.



- 1 a** A gold nanoparticle is 83 nm in diameter. Calculate its diameter in metres, m.



- b** The diameter of a methane molecule is 0.41 nm. Estimate how many times larger the gold nanoparticle is compared to a methane molecule.

## Nanoparticulate materials

Nanoparticles occur naturally. However, modern applications involve manufactured nanoparticles. Substances that consist of nanoparticles are described as being **nanoparticulate**. The uses of these materials depend on the small size of the nanoparticles, and their large **surface area to volume ratios**.

## Did you know?

Smaller gold nanoparticles suspended in water give a red colour, while larger ones give a purple colour.

**Worked example W2**

A cube-shaped nanoparticle has sides of 20 nm.

- a** Calculate its total surface area.

$$\text{surface area} = 6 \times 20 \times 20 = 2400 \text{ nm}^2$$

- b** Calculate its volume.

$$\text{volume} = 20 \times 20 \times 20 = 8000 \text{ nm}^3$$

- c** Calculate its surface area to volume ratio.

$$\text{surface area to volume ratio} = \frac{2400}{8000} = 0.3$$

- 2** A cube-shaped nanoparticle has sides of 2 nm. Calculate its:



- a** total surface area



- b** volume



- c** surface area to volume ratio.



- d** Compare your answer to part **c** with answer **c** in Worked example W2. What happens to the surface area to volume ratio as nanoparticles become smaller?

**Uses of nanoparticulate materials**

Titanium dioxide in bulk is a white solid. It absorbs harmful ultraviolet radiation present in sunlight, which may cause skin cancer. Nanoparticulate titanium dioxide is transparent because its particles are very small, but it still absorbs ultraviolet radiation. This makes it useful for almost-invisible sunscreens.



- 3** Suggest an explanation for why nanoparticles are used in sunscreens.

The very large surface area to volume ratios of nanoparticles makes them useful as catalysts. Stain-resistant clothes treated with nanoparticulate materials stay clean because the nanoparticles catalyse the breakdown of dirt.



- 4** Suggest two reasons that explain why titanium dioxide nanoparticles are used to provide a self-cleaning coating for windows.

**Risks of nanoparticles**

Some scientists are worried that nanoparticulate materials may pose hazards to human health and to the environment. The small size of nanoparticles allows them to be breathed in, or to pass through cell-surface membranes. Their large surface area to volume ratios may allow them to catalyse harmful reactions, or to carry toxic substances bound to their surfaces. The risks are difficult to determine because modern nanoparticulate materials have not been in use for long.



- 5** Some socks are treated with antibacterial silver nanoparticles, stopping the socks from becoming smelly. Describe a possible environmental hazard caused by these nanoparticles.



**C** The cotton fibres in this electron microscope image are coated with titanium dioxide nanoparticles, which make them self-cleaning.

**Checkpoint**

How confidently can you answer the Progression questions?

**Strengthen**

- S1** What are nanoparticulate materials?  
**S2** State some uses and possible hazards of nanoparticles.

**Extend**

- E1** Evaluate the use of nanoparticles in sunscreens.

**Exam-style question**

Give two reasons that explain why some scientists are worried about the possible effects of nanoparticulate materials on humans. (2 marks)



## Qualitative analysis

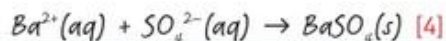
A school technician finds some white crystals in an unlabelled beaker in the laboratory.

The substance in the beaker could be sodium bromide, sodium sulfate, potassium bromide or potassium sulfate.

Explain how the technician could use chemical tests to determine which of the four compounds was left in the beaker. You may include equations in your answer. (6 marks)

### Student answer

The technician can use flame tests to find out whether the substance contains sodium ions or potassium ions [1]. A small sample of the substance is held at the edge of a hot Bunsen burner flame using a clean flame test loop. Sodium ions will produce a yellow flame colour and potassium ions will produce a lilac flame colour [2]. The technician will then need to find out whether the substance contains bromide ions or sulfate ions. A small sample of the substance is dissolved in distilled water and divided in two [3]. A few drops of dilute hydrochloric acid are added (to remove carbonate ions that would give a precipitate), followed by dilute barium chloride solution. A white precipitate of barium sulfate forms if the substance contains sulfate ions:



A few drops of dilute nitric acid are added, followed by dilute silver nitrate solution. A cream precipitate of silver bromide forms if the substance contains bromide ions.

[1] Flame tests must be used because  $\text{Na}^+$  ions and  $\text{K}^+$  ions do not form precipitates with dilute sodium hydroxide solution (unlike  $\text{Ca}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$  ions).

[2] The correct flame colours for the two metal ions are given.

[3] The answer might equally well have started with the tests to identify sulfate ions and bromide ions, rather than with the flame tests.

[4] The answer includes a balanced ionic equation with appropriate state symbols, but the written explanation is already very clear.

### Verdict

This is a strong answer. It shows good knowledge and understanding of how to carry out tests to identify ions in compounds, and the results expected. The answer explains the steps needed in a logical order and includes practical details such as the need to use distilled water in a chemical analysis.

#### Exam tip

Make sure you know how to carry out tests for ions and the observations expected.

# Glossary

<b>absorb</b>	To soak up or take in.	<b>batch process</b>	A method in which substances are made in discrete stages rather than constantly.
<b>acid</b>	A solution that reacts with alkalis, turns litmus red and has a pH of less than 7.	<b>biodegradable</b>	A substance that can be broken down by microorganisms.
<b>acid rain</b>	Rainwater that is more acidic than usual due to air pollution, usually caused by sulfur dioxide and nitrogen oxides.	<b>bioleaching</b>	Using bacteria to extract metals from their ores.
<b>acidic</b>	Containing or having the properties of an acid.	<b>bleach</b>	To bleach means to take the colour out of something.
<b>acidity</b>	The amount of acid in a solution.	<b>boiling point</b>	The temperature at which a substance changes from a liquid to a gas.
<b>activation energy</b>	The minimum amount of energy needed to start a reaction.	<b>bond</b>	The force of attraction between atoms or ions holding them together.
<b>active site</b>	The space in an enzyme where the substrate fits during an enzyme-catalysed reaction.	<b>bond energy</b>	The energy needed to break one mole of a specified covalent bond. It is measured in $\text{kJ mol}^{-1}$ .
<b>actual yield</b>	The actual amount of product obtained from a chemical reaction.	<b>bulk</b>	A substance in the form of lumps or powders is described as being in bulk.
<b>addition polymerisation</b>	A type of polymerisation in which the monomers add on to each other and no small molecules are eliminated.	<b>burette</b>	A piece of apparatus used to accurately measure the volume of solution that has been added during a titration.
<b>addition reaction</b>	A reaction in which reactants combine to form one larger product molecule and no other products.	<b>by-product</b>	Substance produced in a chemical reaction in addition to the desired product.
<b>alcohol</b>	An homologous series of compounds that contain the $-\text{OH}$ functional group.	<b>calibrated</b>	Marked with a scale for accurate readings.
<b>alkali</b>	A solution which contains excess $\text{OH}^-$ ions, turns litmus blue and has a pH greater than 7.	<b>calibration curve</b>	A graph used to determine the concentration of a substance in a sample.
<b>alkali metals</b>	A group of very reactive metals found in group 1 of the periodic table.	<b>carbohydrate</b>	A group of compounds made of carbon, hydrogen and oxygen. Sugars and starch are examples of carbohydrates.
<b>alkaline</b>	Containing or having the properties of an alkaline.	<b>carbon monoxide</b>	A poisonous gas produced from carbon burning without enough oxygen.
<b>alkalinity</b>	The amount of alkali in a solution.	<b>carboxylic acid</b>	An homologous series of compounds that contain the $-\text{COOH}$ functional group.
<b>alkane</b>	A hydrocarbon in which all the bonds between the carbon atoms are single bonds.	<b>catalyst</b>	A substance that speeds up the rate of a reaction without itself being used up.
<b>alkene</b>	A hydrocarbon in which there are one or more double bonds between carbon atoms.	<b>cathode</b>	Negative electrode.
<b>allotrope</b>	A different structural form of an element, e.g. graphite and diamond are allotropes of carbon.	<b>cation</b>	A positively-charged ion formed by losing electrons.
<b>alloy</b>	A metal with one or more other elements (usually metals) added to improve its properties.	<b>causal link</b>	When one thing can be shown to be causing another thing.
<b>alloy steel</b>	Iron with other elements deliberately added to it to make an alloy.	<b>ceramic</b>	A hard, durable, non-metallic material which is generally unaffected by heat, e.g. china and glass.
<b>amino acid</b>	A group of compounds that are the monomers used to form proteins.	<b>chemical analysis</b>	Using chemical reactions or sensitive machines to identify and measure substances in a sample.
<b>anaerobic respiration</b>	A type of respiration that does not need oxygen.	<b>chemical cell</b>	A device that produces a voltage due to reactions between reactants stored inside it, until one of the reactants is used up.
<b>anion</b>	A negatively-charged ion formed by gaining electrons (usually a non-metal ion).	<b>chemical property</b>	How a substance reacts with other substances.
<b>anode</b>	Positive electrode.	<b>chlorination</b>	The process of adding chlorine to a substance, often to water.
<b>aqueous solution</b>	A mixture that is formed when a substance is dissolved in water.	<b>chromatogram</b>	The piece of paper showing the results of chromatography.
<b>aquifer</b>	An underground layer of rock containing groundwater which can be extracted using a well or pump.	<b>chromatography</b>	A technique for separating the components of a mixture, e.g. different food colouring agents.
<b>atmosphere</b>	Layer of gases that surrounds the Earth.	<b>clay ceramic</b>	A hard, durable material including brick, porcelain and china.
<b>atom</b>	The smallest neutral part of an element that can take part in chemical reactions.	<b>climate change</b>	Changes that happen to the global weather patterns as a result of global warming.
<b>atom economy</b>	The percentage, by mass, of reactants that are converted into useful products.	<b>closed system</b>	When substances cannot enter or leave an observed environment, e.g. a stoppered test tube.
<b>atomic number</b>	The number of protons in the nucleus of an atom (symbol $Z$ ). It is also known as the proton number.	<b>combustion</b>	A chemical reaction in which a compound reacts with oxygen.
<b>attractive forces</b>	The weak forces of attraction between molecules.	<b>complete combustion</b>	Combustion of hydrocarbons with enough oxygen present to convert all the fuel into carbon dioxide and water.
<b>Avogadro constant</b>	This is the number of particles in one mole of a substance ( $6.02 \times 10^{23} \text{ mol}^{-1}$ ).	<b>composite material</b>	A mixture of two or more materials with contrasting properties, combined to produce a material with properties of both.
<b>Avogadro's law</b>	If the temperature and pressure are the same, equal volumes of different gases contain an equal number of molecules.	<b>composition</b>	The substances and their proportions that make something up. For example, the composition of today's atmosphere is 78% nitrogen, 21% oxygen, 1% argon and other gases.
<b>balanced equation</b>	Description of a reaction using the symbols and formulae of the reactants and products, so that the number of 'units' of each element to the left of the arrow is the same as those to the right of the arrow.	<b>compound</b>	A substance that can be split into simpler substances because it contains the atoms of two or more elements joined together.
<b>base</b>	A substance that will react with an acid to form only salt and water.	<b>compressive strength</b>	A measure of how well a substance resists squashing.
		<b>concentrated</b>	Containing a large amount of solute dissolved in a small volume of solvent.



<b>concentration</b>	The amount of a solute dissolved in a certain volume of solvent.	<b>electrode</b>	A rod made of a metal or graphite that carries the current into or out of the electrolyte.
<b>condensation polymerisation</b>	When monomers join together and eliminate a small molecule, such as water.	<b>electrolysis</b>	The process in which energy transferred by a direct electrical current decomposes electrolytes.
<b>condense</b>	When a gas turns in to a liquid.	<b>electrolyte</b>	An ionic compound that is molten or dissolved in water.
<b>confirmatory test</b>	A chemical test carried out to check the conclusion from the results of another test.	<b>electron</b>	A tiny particle with a negative charge and very little mass.
<b>continuous process</b>	A method of manufacture in which substances are made constantly.	<b>electron shell</b>	Areas around a nucleus that can be occupied by electrons and are usually drawn as circles. Also called an electron energy level or an 'orbit'.
<b>correlation</b>	A relationship between two variables, so that if one variable changes so does the other. This can be positive or negative.	<b>electronic configuration</b>	The arrangement of electrons in shells around the nucleus of an atom.
<b>corrosion</b>	The gradual deterioration of a substance when it reacts with substances in the environment, for example when a metal oxidises in air.	<b>electroplating</b>	Using electricity to coat one metal with a thin layer of another metal.
<b>covalent bond</b>	The bond formed when a pair of electrons is shared between two atoms.	<b>electrostatic force</b>	The force of attraction between oppositely-charged particles, and force of repulsion between particles with the same charge.
<b>covalent, giant molecular structure</b>	Three-dimensional lattice of carbon atoms linked by covalent bonds.	<b>element</b>	A simple substance made up of only one type of atom.
<b>covalent, simple molecular structure</b>	Two or more atoms covalently bonded together to form a distinct unit.	<b>emission spectra</b>	A set of wavelengths of light or electromagnetic radiation showing which wavelengths have been given out (emitted) by a substance.
<b>cracking</b>	A chemical reaction in which large alkane molecules are split into two or more smaller alkanes and alkenes.	<b>emit</b>	To give out.
<b>crude oil</b>	A mixture of hydrocarbons formed from dead microscopic organisms by heat and pressure over millions of years.	<b>empirical formula</b>	The formula showing the simplest whole number ratio of atoms of each element in a compound.
<b>crystallisation</b>	Separating the solute from a solution by evaporating the solvent.	<b>endothermic</b>	A type of reaction in which energy from the surroundings is transferred to the products, e.g. photosynthesis.
<b>crystals</b>	Solids that are made up of a regular, repeated pattern of atoms, molecules or ions which form fixed shapes with flat surfaces and sharp edges.	<b>end-point</b>	When just enough solution has been added from the burette to react with all the solution in the flask in a titration experiment.
<b>delocalised electron</b>	An electron that is free to move and can carry an electrical current.	<b>enzyme</b>	A protein produced by living organisms that acts as a catalyst to speed up the rate of a reaction.
<b>denature</b>	A denatured enzyme is one where the shape of the active site has changed so much that its substrate no longer fits and the reaction can no longer happen.	<b>ester link</b>	This link is present in all polyester molecules. It consists of $-\text{COO}-$ .
<b>desalination</b>	A process that produces fresh drinking water by separating the water from the salts in salty water.	<b>ethanoic acid</b>	The carboxylic acid which contains two carbon atoms and is the main acid in vinegar.
<b>desiccant</b>	A substance that absorbs water or vapour.	<b>evaporate</b>	When a liquid turns in to a gas.
<b>diatomic</b>	Two atoms chemically bonded together.	<b>exothermic</b>	A type of reaction in which energy is transferred to the surroundings from the reactants, e.g. combustion.
<b>dilute</b>	A low concentration of a solute in a solution.	<b>extraction</b>	A process in which a metal is obtained from its ore.
<b>discharged</b>	In electrolysis, an ion is discharged when it gains or loses electrons to form an element.	<b>feedstock</b>	Raw material. A substance used to make other substances.
<b>disinfectant</b>	Something that destroys or neutralises disease-carrying microorganisms.	<b>fermentation</b>	Anaerobic respiration occurring in microorganisms.
<b>displacement reaction</b>	When a more reactive element displaces a less reactive element from one of its compounds.	<b>fertiliser</b>	Soluble compounds added to soil to replace the minerals used up by plants.
<b>dissociate</b>	Breaking up of a compound in to simpler components.	<b>filtrate</b>	A solution that has passed through a filter.
<b>distillate</b>	The liquid produced by condensing gases during distillation.	<b>filtration</b>	Using a filter to separate insoluble substances from a liquid.
<b>distillation</b>	The process of separating a liquid from a mixture by evaporating the liquid and then condensing it (so that it can be collected).	<b>finite resource</b>	Something useful that is no longer made or which is being made very slowly.
<b>dot and cross diagram</b>	A diagram to explain what happens when a bond is formed. It uses dots and crosses to represent the electrons of different atoms.	<b>flame photometer</b>	A machine used to identify metal ions in solution and to determine their concentration.
<b>double bond</b>	The covalent bond formed when two pairs of electrons are shared between the same two atoms.	<b>fossil fuel</b>	A fuel formed from the dead remains of organisms over millions of years, i.e. coal, oil and natural gas.
<b>ductile</b>	A substance that can be stretched out to make a thin wire.	<b>fraction</b>	A component of a mixture that has been separated by fractional distillation.
<b>dynamic equilibrium</b>	When the forwards and backwards reactions in a reversible chemical reaction are occurring at the same rate.	<b>fractional distillation</b>	A method of separating a mixture of liquids with different boiling points into individual components (fractions).
<b>effervescence</b>	The formation of gas bubbles in a liquid due to a chemical reaction occurring.	<b>fractionating column</b>	A long column used for fractional distillation. It is warmer at the bottom than at the top.
<b>electrical conductivity</b>	Allowing electricity to pass through.	<b>fuel cell</b>	A device that produces a voltage due to reactions involving a fuel and oxygen, for as long these reactants are supplied.
		<b>fullerene</b>	A molecule in which each carbon atom is covalently bonded to three other carbon atoms, forming spheres or tube shapes.
		<b>functional group</b>	An atom or group of atoms in a molecule that is mainly responsible for the molecule's chemical reactions and properties.



<b>galvanising</b>	Coating iron or steel with a thin layer of zinc to improve its resistance to rusting.	<b>limiting reactant</b>	The reactant that determines the amount of product formed in a chemical reaction. Any other reactants will be present in excess.
<b>general formula</b>	The formula showing the proportions of different atoms in molecules of an homologous series. For example, alkenes have the general formula $C_nH_{2n}$ .	<b>lubricant</b>	A substance placed between two moving surfaces to reduce the friction between them.
<b>glass</b>	A solid produced by cooling molten substances. The atoms are joined to form a giant structure without crystals.	<b>malleable</b>	A substance that can be hammered or rolled into shape without shattering.
<b>global warming</b>	The increase in the Earth's average temperature likely to be caused by increased amounts of carbon dioxide in the atmosphere.	<b>mass number</b>	The total number of protons and neutrons in the nucleus of an atom (symbol A). It is also known as the nucleon number.
<b>graphene</b>	An allotrope of carbon consisting of a sheet that is one atom thick, with atoms arranged in a honeycomb shape.	<b>matrix</b>	In a composite material, it is the substance that binds the reinforcement material together.
<b>greenhouse effect</b>	When gases in the atmosphere absorb energy transferred by infrared waves from the Earth, which causes the atmosphere to be warmer than it otherwise would be.	<b>mean</b>	An average calculated by adding up the values of a set of measurements and dividing by the number of measurements in the set.
<b>greenhouse gas</b>	A gas that helps to trap 'heat' in the atmosphere. Carbon dioxide, methane and water vapour are greenhouse gases.	<b>melting point</b>	The temperature at which a substance changes from the solid state to the liquid state when heated, or from the liquid state to the solid state when cooled.
<b>group</b>	A vertical column of elements in the periodic table. Elements in the same group generally have similar properties.	<b>metal</b>	Any element that is shiny when polished, conducts heat and electricity well, is malleable and flexible and often has a high melting point.
<b>haemoglobin</b>	The red, iron-containing pigment found in red blood cells.	<b>metallic bonding</b>	The type of bonding found in metals. You can think of it as positively-charged ions in a 'sea' of negatively-charged electrons.
<b>half equation</b>	A chemical equation written to describe an oxidation or reduction half-reaction.	<b>mixture</b>	A substance containing two or more different substances that are not joined together.
<b>halide</b>	A compound formed between a halogen and another element such as a metal or hydrogen.	<b>mobile phase</b>	In paper chromatography, this is when the solvent moves along the paper carrying the dissolved samples with it.
<b>halide ion</b>	A negatively-charged ion formed from one of the group 7 elements.	<b>molar gas volume</b>	The volume occupied by one mole of molecule of any gas. It is $24\text{ dm}^3$ or $24\,000\text{ cm}^3$ at room temperature and pressure.
<b>halogen</b>	An element in group 7 of the periodic table.	<b>mole</b>	A mole of something is $6 \times 10^{23}$ of it. The mass of a mole of a substance is the relative formula mass expressed in grams.
<b>hazard</b>	Something that could cause harm.	<b>mole ratio</b>	The ratio in moles between substances in a balanced equation. It is the balancing numbers of the substances.
<b>homologous series</b>	A family of compounds that have the same general formula and similar properties, but have different numbers of carbon atoms.	<b>molecular formula</b>	The formula showing the actual number of atoms of each element in a molecule of a compound.
<b>hydrocarbon</b>	A compound containing only hydrogen and carbon atoms.	<b>molecular</b>	Refers to substances that are made up of molecules.
<b>ignite</b>	To start burning.	<b>molecule</b>	A particle consisting of two or more atoms joined together by covalent bonding.
<b>impurity</b>	Unwanted substance found mixed into a useful substance.	<b>monomer</b>	A small molecule that can join with other molecules like itself to form a polymer.
<b>incinerated</b>	This occurs when a substance is burned.	<b>nanoparticle</b>	Piece of a material consisting of a few hundred atoms, and between 1 nm and 100 nm in size.
<b>incomplete combustion</b>	When a substance reacts only partially with oxygen, such as when carbon burns in air producing carbon dioxide, carbon monoxide and soot (unburnt carbon).	<b>nanoparticulate</b>	A material consisting of nanoparticles is described as being nanoparticulate.
<b>indicator</b>	A substance which can change colour depending on the pH of a solution.	<b>native state</b>	The native state of an element is when it is not combined with other elements in compounds.
<b>inert</b>	Does not react.	<b>natural gas</b>	A fossil fuel formed from the remains of microscopic dead plants and animals that lived in the sea.
<b>infrared</b>	Electromagnetic radiation that we can feel as heat.	<b>naturally occurring polymer</b>	A substance that exists naturally as a polymer in plants, animals etc., such as DNA, starch and proteins.
<b>insoluble</b>	A substance that cannot be dissolved in a certain liquid.	<b>neutral</b>	A liquid that is neither acidic nor alkaline and has a pH of 7.
<b>intermolecular force</b>	A weak force of attraction between molecules.	<b>neutralisation</b>	A reaction in which an acid reacts with a base to produce a salt and water only.
<b>ion</b>	An atom or group of atoms with an electrical charge due to the gain or loss of electrons.	<b>neutralise</b>	To make a solution neither acidic nor alkaline. During neutralisation, a base reacts with an acid, forming a salt and water.
<b>ionic bond</b>	A strong electrostatic force of attraction between oppositely-charged ions.	<b>neutron</b>	A particle found in the nucleus of an atom having zero charge and mass of 1 (relative to a proton).
<b>ionic compound</b>	A substance made up of ions of different elements.	<b>nitrogenous</b>	Containing or supplying nitrogen, for example in a fertiliser.
<b>ionic equation</b>	A balanced equation that only shows the ions that react together. The spectator ions are not included in the equation.	<b>noble gas</b>	An unreactive gas in group 0 of the periodic table.
<b>isomer</b>	Molecules with the same molecular formula but different arrangements of atoms are called isomers.	<b>non-enclosed system</b>	Another term for an open system.
<b>isotope</b>	Atoms of an element with the same number of protons (atomic number) but different mass numbers due to different numbers of neutrons.	<b>non-metal</b>	An element that is not shiny and does not conduct heat or electricity well.
<b>lattice structure</b>	An arrangement of many particles that are bonded together in a fixed, regular, grid-like pattern.		
<b>law of conservation of mass</b>	The idea that mass is never lost or gained during a chemical reaction or physical change.		
<b>leachate</b>	A solution produced when water or another solvent passes through a mixture of substances and dissolves some of them.		



<b>non-renewable</b>	Any energy resource that will run out because you cannot renew your supply of it, e.g. oil.	<b>precipitation reaction</b>	A reaction in which an insoluble product is formed from two soluble reactants.
<b>nuclear fission</b>	When the nucleus of a large atom such as uranium, splits into two smaller nuclei.	<b>prediction</b>	What you think will happen in an experiment (usually given with a reason of why you think this).
<b>nucleotide</b>	The monomers that make up nucleic acids such as DNA.	<b>product</b>	A substance formed in a reaction.
<b>nucleus</b>	The central part of an atom or ion.	<b>protein</b>	A polymer made up of amino acids.
<b>opaque</b>	Material that does not let light through. It is not possible to see through an opaque substance.	<b>proton</b>	A particle found in the nucleus of an atom, having a positive charge and the same mass as a neutron.
<b>open system</b>	A system into or from which substances can enter or leave, such as a reaction inside an open test tube.	<b>pure</b>	A single substance with a fixed composition that does not have anything else mixed with it.
<b>organic compound</b>	A compound that has a central framework of carbon atoms onto which hydrogen and other atoms are attached. Methane ( $\text{CH}_4$ ) is organic but carbon dioxide is not (because it contains no hydrogen atoms).	<b>rate</b>	How quickly something happens.
<b>outer electron shell</b>	The electron shell (or energy level) that is furthest away from a nucleus but which still contains one or more electrons.	<b>reactant</b>	A substance used up in a chemical reaction.
<b>oxidation</b>	A reaction in which oxygen is added to a chemical; loss of electrons by an atom or negative ion.	<b>reaction pathway</b>	A series of reactions needed to make a particular product.
<b>oxide of nitrogen</b>	Any one of a variety of gaseous compounds consisting of only nitrogen and oxygen atoms. Together they are often represented as $\text{NO}_x$ .	<b>reaction profile</b>	A diagram to show how the energy stored in a substance changed during the course of a chemical reaction.
<b>oxidising agent</b>	A substance that causes another substance to be oxidised in an oxidation reaction.	<b>reactivity series</b>	A list of metals in order of reactivity with the most reactive at the top.
<b>paper chromatography</b>	Chromatography carried out by spotting drops of the same samples onto paper and then allowing a solvent to move up the paper. Different components in the samples travel up the paper in the solvent at different rates.	<b>recycling</b>	Converting waste materials into new products.
<b>particle</b>	A tiny piece of matter that everything is made out of.	<b>red blood cell</b>	A biconcave disc containing haemoglobin that gives blood its red colour and carries oxygen around the body to the tissues. Also known as an erythrocyte.
<b>particle model</b>	Another term for kinetic theory.	<b>redox reaction</b>	A reaction in which both oxidation and reduction occur.
<b>percentage yield</b>	The actual yield divided by the theoretical yield, as a percentage.	<b>reduction</b>	A reaction in which a substance loses oxygen or gains electrons.
<b>period</b>	A horizontal row in the periodic table.	<b>reinforcement</b>	In a composite material, the substance that is bound together by the matrix material.
<b>periodic table</b>	The chart in which the elements are arranged in order of increasing atomic number.	<b>relative atomic mass (RAM, <math>A_r</math>)</b>	The mean mass of an atom relative to the mass of an atom of carbon-12, which is assigned a mass of 12. The RAM of an element is the mean relative mass of the isotopes in the element.
<b>petrochemical</b>	A substance made from crude oil.	<b>relative charge</b>	The electrical charge on something compared with something else. The electrical charge used for the comparison is often given the value of 1.
<b>pH scale</b>	A scale going up to 14 showing acidity or alkalinity. Numbers below 7 are acids; numbers above 7 are alkalis; pH 7 is neutral.	<b>relative formula mass (<math>M_r</math>)</b>	The sum of the relative atomic masses of all the atoms in a formula.
<b>photosynthesis</b>	A series of enzyme-catalysed reactions carried out in the green parts of plants. Carbon dioxide and water combine to form glucose. This process requires light energy from sunlight.	<b>relative mass</b>	The mass of something compared to the mass of something else, which is often given the mass of 1.
<b>physical change</b>	A change in which no new substances are formed, such as changes of state.	<b>renewable source</b>	Source of raw materials that will not run out.
<b>physical property</b>	A description of how a material behaves and responds to forces and energy. For example, hardness is a physical property.	<b>repeating unit</b>	The part of a polymer that can be repeated many times to form the polymer chain.
<b>phytoextraction</b>	Using plants to extract metals from their ores.	<b>residue</b>	Material remaining in the filter after a mixture has passed through it.
<b>pipette</b>	A piece of apparatus which can be used in a titration to accurately measure a set volume of a solution.	<b>resolution</b>	The smallest change that can be measured by an instrument. For example, in a microscope it is the smallest distance between two points that can be seen as two points and not blurred into one point.
<b>plasticiser</b>	A substance added to a polymer during its manufacture to make the polymer softer and more flexible.	<b>reversible reaction</b>	A chemical reaction that can work in both directions.
<b>pollutant</b>	A substance that harms living organisms when released into the environment.	<b><math>R_f</math> value</b>	The ratio of the distance travelled by a solute on a chromatogram to the distance travelled by the solvent under the same conditions.
<b>polyatomic ions</b>	A group of atoms that have a positive or negative charge due to the loss or gain of electrons, e.g. nitrate $\text{NO}_3^-$ .	<b>risk</b>	The chance of a hazard causing harm.
<b>polyester</b>	This is a polymer that contains large numbers of ester links.	<b>risk assessment</b>	Identification of the hazards of doing an experiment and ways of reducing the risk of harm from those hazards.
<b>poly(ethene)</b>	A common polymer made of ethene monomers.	<b>rusting</b>	The corrosion of iron or steel (water and oxygen must be present for rusting to occur).
<b>polymer</b>	A long-chain molecule made by joining many smaller molecules (monomers) together.	<b>sacrificial protection</b>	Using a more reactive metal to protect iron from rusting.
<b>polymerisation</b>	A reaction in which a large number of small molecules (monomers) join together to form a long chain molecule (polymer).	<b>salt</b>	A compound formed by neutralisation of an acid by a base.
<b>precipitate</b>	An insoluble substance that is formed when two soluble substances react together in solution.	<b>saturated</b>	A molecule that contains only single bonds between the carbon atoms in a chain.
<b>precipitation</b>	A reaction in which a precipitate is formed.	<b>saturated solution</b>	Contains the maximum amount of solute that can dissolve in that amount of solvent at that temperature.
		<b>sedimentation</b>	The process in which rock grains and insoluble substances sink to the bottom of a liquid.

<b>side reaction</b>	A reaction which takes place at the same time as another main reaction.	<b>tarnish</b>	A thin layer that forms on a metal due to oxidation. A metal is also said to tarnish as this layer forms.
<b>simple distillation</b>	The process of separating a liquid from a mixture by evaporating the liquid and then condensing it so that it can be collected.	<b>tensile strength</b>	A measure of how well a substance resists stretching.
<b>solute</b>	Describes a substance that dissolves in a liquid to make a solution.	<b>theoretical yield</b>	The maximum calculated amount of a product that could be formed from a given amount of reactants.
<b>solution</b>	Formed when a substance has dissolved in a liquid.	<b>titration</b>	A technique in volumetric analysis that is used to find the exact volumes of solutions which react with each other.
<b>solvent</b>	Describes the liquid in which a substance dissolves to make a solution.	<b>toxic</b>	Poisonous.
<b>spectator ions</b>	These are ions that do not change during a reaction.	<b>transition metal</b>	A metal element placed in the block between groups 2 and 3 in the periodic table.
<b>spectrum</b>	Individual components of light arranged in order of wavelength or frequency.	<b>transparent</b>	A coloured or colourless material that light can travel through without scattering.
<b>stainless steel</b>	Alloy steel containing elements such as chromium, to resist rusting.	<b>universal indicator</b>	A mixture of different indicators giving a different colour at different points on the pH scale.
<b>standard solution</b>	A solution containing a known concentration of a substance.	<b>unsaturated</b>	A molecule that contains one or more double bonds between carbon atoms in a chain.
<b>starch</b>	A polymer carbohydrate that is made by the joining together of glucose molecules.	<b>valency</b>	The number of covalent bonds formed by an atom, or the charge number of the ion formed by an atom.
<b>state of matter</b>	One of three different forms that a substance can have: solid, liquid or gas.	<b>variable</b>	A factor that can change.
<b>state symbol</b>	A letter or letters to show the state of a substance.	<b>viscosity</b>	How thick or runny a liquid is. Low viscosity is very runny; high viscosity is thick.
<b>stationary phase</b>	The surface through which the solvent and dissolved substances move in chromatography.	<b>volcanic activity</b>	The release of gases and/or molten rock by volcanoes.
<b>still</b>	A piece of apparatus used to carry out distillation or fractional distillation.	<b>volumetric flask</b>	A flask which is accurately calibrated to hold a given volume of solution.
<b>stoichiometry</b>	The molar ratio of the reactants and products in a chemical reaction.	<b>weak acid</b>	An acidic solute that does not dissociate completely into ions when it dissolves.
<b>strong acid</b>	An acidic solute that dissociates completely into ions when it dissolves.	<b>weathering</b>	When rocks are broken up by physical, chemical or biological processes.
<b>structural formula</b>	The formula showing the symbols for each atom in a compound with straight lines joining them to represent the covalent bonds.	<b>X-ray</b>	Electromagnetic radiation that has a shorter wavelength than UV but longer than gamma rays.
<b>subatomic particle</b>	A particle that is smaller than an atom, such as a proton, neutron or electron.	<b>yield</b>	The amount of useful product that you can get from something.
<b>substrate</b>	A substance that is changed during a reaction.		
<b>sugar</b>	Soluble carbohydrate made up of small molecules, e.g. glucose formula $C_6H_{12}O_6$ .		
<b>surface area : volume ratio (SA : V)</b>	The total amount of surface area of an object divided by its volume.		
<b>synthetic polymer</b>	A polymer that is manufactured in a laboratory or factory.		



## A

A, (relative atomic mass) 23  
 absorption 166  
 acid rain 158  
 acidic solutions 52  
 acidity 52  
 acids 52–53, 54–55  
     reaction with metals 66  
     strong 55  
     weak 55  
 activation energy 138, 146  
     catalysts 142–143  
 active sites 143  
 actual yield 108  
 addition polymerisation 184–185  
 addition reactions 175  
 alcohols 178–179  
     ethanol 176–177  
 alkali metals 128–129  
 alkaline solutions 52  
 alkalinity 52  
 alkalis 52–53, 60–61, 64–65  
 alkanes 154, 172, 178  
     homologous series 154–155  
     reactions 174–175  
 alkenes 161, 173  
     reactions 174–175  
 allotropes of carbon 44–45  
 alloy steels 102  
 alloys 102–103, 203  
     uses 104–105  
 amino acids 185  
 ammonium ions, tests for 197  
 ammonium nitrate fertiliser 120  
 ammonium sulfate 120–121  
 anaerobic respiration 177  
 anions 34, 80, 198  
 anode 80, 100  
 aqueous solutions  
     acids and alkalis 52  
     ionic compounds 39  
 aquifers 15  
 atmosphere  
     changing 164–165  
     composition 162  
     early 162–163  
     present-day 166–167  
 atom economy 110–111  
 atomic number 20  
     periodic table 28–29  
 atomic structure 18–19  
 atoms 2, 18  
 attractive forces 3  
 Avogadro constant 76  
 Avogadro's law 118

## B

balanced equations 61  
 bases 56  
 batch processes 121  
 biodegradable 190  
 bioleaching 89  
 bleaches 131  
 boiling point 3  
     ionic compounds 38  
     molecular compounds 42  
 bond energy 147  
 bonding 42  
     covalent 40–41, 42, 48  
     giant covalent 48

ionic 34–35, 48  
 metallic 46, 48  
 models 48–49  
 bonds  
     breaking and making 146  
     covalent 147  
     energy calculations 147  
 bromine water test 175  
 bulk materials 206  
 by-products 110, 123, 125

## C

calibration 112  
 calibration curves 195  
 carbohydrates 176  
 carbon allotropes 44–45  
 carbon monoxide 157  
 carbonates  
     reactions with acids 67  
     test for 198  
 carboxylic acids 182–183  
 catalysts 142–143  
     cracking 160  
 cathode 80, 100  
 cations 34, 80, 194  
 causal links 166  
 ceramics 202  
 chemical cells 124  
 chemical properties of substances 3  
     elements 26  
 chlorination 15  
 chromatogram 8  
 chromatography 8  
     paper 8–9  
 climate change 166–167, 168–169  
     effects 168–169  
     limiting the impact 169  
 closed systems 74, 94, 122  
 combustion 156–157, 174  
 complete combustion 156, 174  
 composite materials 204–205  
 compounds 4  
     ionic 36–39  
     molecular 42–43  
 compressive strength 205  
 concentrated solutions 54  
 concentration 53, 112–113  
 condensation 10, 152  
 condensation polymerisation 188–189  
 conservation of mass, law of 74–75  
 continuous processes 121  
 correlations 166  
 corrosion 91, 98–99  
 covalent bonds 40–41, 42, 147  
     giant molecular structures 45  
     simple molecular structures 42  
 cracking 160, 190  
 crude oil 150  
     fractional distillation 152–153  
 crystallisation 6, 57  
     laboratory practice 7  
 crystals 36

## D

denaturing of enzymes 143  
 desalination 14  
 desiccants 99  
 diatomic molecules 130  
 dilute solutions 54

discharged ions 84  
 disinfectants 131  
 displacement reactions 87, 132, 145  
 dissociation 55  
 distillate 177  
 distillation  
     fractional 11, 152–153  
     simple 14  
     water 10–11, 14  
 dot and cross diagrams 40  
 double bonds 40  
 drinking water 14–15  
 ductile substances 96, 104  
 dynamic equilibrium 94–95  
     factors affecting 122–123

## E

effervescence 66  
 electrical conductivity  
     ionic compounds 38–39  
     metals 47  
     molecular compounds 43  
 electrodes 80  
     reactions at 81  
 electrolysis 80–81  
     products 84–85  
     salt solutions 84–85  
 electrolytes 80, 100  
 electron shells 18, 30  
     outer 40  
 electronic configurations 30–31  
     noble gases 34  
 electrons 18  
 electroplating 100–101  
 electrostatic forces 34  
 elements 4, 18, 42  
     groups 29  
     periodic table 26–27  
 emission 166  
 emission spectra 195  
 empirical formulae 73  
 endothermic reactions 138, 144–145  
     equilibrium 95  
 end-point of a titration 65  
 energy changes during reactions 146–147  
 enzymes 143, 176  
     denaturing 143  
 equations  
     balanced 61  
     half equations 66, 81, 87  
     ionic 66  
 equilibrium, dynamic 94–95  
     factors affecting 122–123  
 ester links 188  
 ethanoic acid 182  
 ethanol 176–177  
 evaporation 10, 152  
 exam preparation examples  
     alcohols and carboxylic acid 192  
     atomic structure and periodic table 32  
     bonding 50  
     electrolysis 106–107  
     empirical formula 78  
     fuels 170  
     isotopes 24  
     magnesium sulfate 70  
     qualitative analysis 208  
     reaction rates 148

substance separation and purification 16  
 exothermic reactions 138, 144–145  
     equilibrium 95  
 extraction of metals 88

## F

feedstock 151  
 fermentation 176–177  
 fertilisers 120–121  
 filtrate 7  
 filtration 6, 57  
     laboratory practice 7  
 finite resources 190  
 flame photometry 195  
 flame tests 194  
 fossil fuels 151  
 fractional distillation 11, 152–153, 177, 190  
 fractionating columns 152  
 fractions (of distillation) 177  
 fuel cells 124–125  
 fuels 151  
     for cars 161  
     pollution 158–159  
 fullerenes 44  
 functional groups 173, 179, 188

## G

galvanising 101  
 gases, molar volumes of 118–119  
 general formulae 155, 172  
 giant structures of carbon 45  
 glass 202  
 global warming 166  
 graphene 44  
 greenhouse effect 166  
 greenhouse gases 125, 161, 166  
 groups (of the periodic table) 29  
     group 0 elements 134–135  
     group 1 elements 128–129  
     group 7 elements 130–131, 132–133  
     valency 41

## H

Haber process 120–121  
 haemoglobin 157  
 half equations 66, 81, 87  
 halide ions 130  
     tests for 199  
 halogens 130–131  
     reactivity 132–133  
 hazards 7  
 homologous series 154, 172, 178  
 hydrocarbons 150–151, 172  
     breaking down 160–161

## I

ignition 153  
 incomplete combustion 157  
 incomplete combustion 174  
 indicators (pH) 52  
 inert electrodes 84  
 inert substances 28  
 infrared radiation 166  
 insoluble substances 6, 69  
 intermolecular forces 42  
 ionic bonds 34–35  
 ionic compounds 36  
     properties 38–39

ionic equations 66  
 ionic formulae 36  
 ionic lattices 36–37  
 ions 53

neutralisation 64–65  
 spectator ions 66  
 tests for negative ions 198–199  
 tests for positive ions 196–197

isomers 173  
 isotopes 22–23

## L

laminates 205  
 lattices

giant 46  
 ionic 36–37

law of conservation of mass 74–75  
 leachate 89  
 life cycle assessment (LCA) 93  
 limiting reactant 76  
 lubricants 45

## M

malleable substances 96, 104, 203  
 malleability of metals 47  
 mass conservation 74–75  
 mass number 21  
 masses 72  
 materials 202–203  
 bulk 206  
 composites 204–205

matrix 204  
 mean 23

melting point 3, 5  
 ionic compounds 38  
 molecular compounds 42

metals 46, 203  
 biological extraction  
 methods 89  
 corrosion 91  
 electrical conductivity 47  
 extraction as reduction 90  
 malleability 47  
 ores 88–89  
 properties 46–47  
 reaction with acids 66  
 reaction with carbonates 67  
 reactivity series 66  
 recycling 92  
 structure and bonding 46  
 uses 104–105

mixtures 4–5  
 separation 10

mobile phase 8  
 molar volume of gases 118–119  
 mole ratio 114  
 molecular compounds 42–43  
 molecular formulae 40, 72, 154  
 working out 41  
 molecular substances 40  
 molecules 2, 40  
 diatomic 130  
 moles 76–77  
 monomers 43, 184, 203

## N

nanoparticles 206–207  
 native state metals 88

natural gas 150, 151  
 naturally-occurring polymers 185  
 negative ions, tests for 198–199  
 neutral solutions 52  
 neutralisation reactions 56, 60, 64–65, 145  
 neutrons 18  
 nitrogen oxides 159  
 nitrogenous fertilisers 120  
 noble gases 134–135  
 non-enclosed systems 74  
 non-metals 46  
 non-renewable materials 151, 190  
 nuclear atom 20  
 nuclear fission 22  
 nucleotides 185  
 nucleus 18

## O

oceans 163, 164  
 oil fractions 152  
 OILRIG mnemonic 66, 81, 133  
 opaque materials 202  
 open systems 94  
 ores 88–89  
 organic compounds 178  
 outer electron shells 40  
 oxidation 66, 81, 87, 90–91, 98, 174, 182  
 OILRIG mnemonic 66, 81, 133  
 oxides of nitrogen 159  
 oxidising agents 182  
 oxygen in the atmosphere 163, 164–165

## P

pair reversals 29  
 paper chromatography 8–9  
 particle model of matter 2  
 particles 2  
 percentage yield 108  
 periodic table 20  
 atomic number 28–29  
 electronic configurations 30–31  
 elements 26–27  
 group 0 elements 134–135  
 group 1 elements 128–129  
 group 7 elements 130–131, 132–133  
 valency 41

periods (of the periodic table) 29  
 petrochemicals 151  
 pH meters 54  
 pH scale 52  
 photometry 195  
 photosynthesis 164  
 physical changes 3  
 physical properties of substances 5  
 elements 26

phytoextraction 89  
 plasticisers 203  
 pollution 158–159  
 polyatomic ions 37  
 polyester 188  
 poly(ethylene) 43  
 polymerisation

addition 184–185  
 condensation 188–189

polymers 43, 184, 203  
 naturally-occurring 185  
 problems with 190–191

properties and uses 186–187  
 synthetic 185

positive ions, tests for 196–197  
 potable (drinkable) water 14–15  
 practicals

acid-base titration 116–117  
 alcohol combustion 180–181  
 copper sulfate electrolysis 82–83  
 copper sulfate preparation 58–59  
 inks 12–13  
 ion identification 200–201  
 neutralisation 62–63  
 reaction rates 140–141

precipitates 14, 68, 74, 196  
 precipitation 68, 145  
 precipitation reactions 196  
 predictions about elements 27  
 products 136

masses of 75

proteins 185  
 protons 18  
 pure substances 4

## R

$R_f$  values 8–9  
 reactants 136  
 limiting 76  
 masses of 75  
 reaction pathways 111, 123  
 reaction profiles 142, 144, 146  
 reaction rates 136–137  
 factors affecting 138–139  
 reactions 76–77  
 displacement 87, 132, 145  
 energy changes 146–147  
 neutralisation 145  
 redox 87, 133  
 reversible 94  
 temperature changes 145

reactivity 86–87  
 reactivity series 66, 86  
 recycling 92–93  
 red blood cells (erythrocytes) 157  
 redox reactions 87, 90, 133  
 reduction 66, 81, 87, 90–91  
 metal extraction 90  
 OILRIG mnemonic 66, 81, 133

reinforcement of materials 204  
 relative atomic masses (RAM) 23, 26, 29  
 relative charges 18  
 relative formula mass 72  
 relative masses 18  
 renewable sources 179  
 repeating units 185  
 residue 7  
 resolution 167  
 reversible reactions 94, 120  
 risk 7  
 risk assessment 7  
 rusting 91, 99

## S

sacrificial protection 99  
 salts 56, 130  
 insoluble 69  
 soluble 57, 65  
 saturated compounds 161  
 saturated solutions 6

sea water purification 14  
 sedimentation 15  
 side reactions 109  
 simple distillation 14  
 solubility 68–69  
 solutes 6, 74  
 solutions 6, 74  
 concentrated 54  
 dilute 54  
 saturated 6  
 solvents 6, 74  
 spectator ions 66, 87  
 spectrum of emitted light 195  
 stainless steels 102  
 standard solutions 195  
 starch 176, 185  
 state changes 3  
 state symbols 56  
 states of matter 2–3  
 stationary phase 8  
 still (distillation equipment) 10  
 stoichiometry 77  
 strong acids 55  
 structural formulae 154  
 subatomic particles 18  
 substances

insoluble 6  
 mixtures 4–5  
 pure 4

substrates 143  
 sugars 176, 185  
 sulfates, test for 198–199  
 sulfur dioxide 158  
 surface area to volume ratios 206  
 synthetic polymers 185

## T

tarnishing 91, 98  
 tensile strength 204  
 theoretical yield 108  
 tin plating 101  
 titrations 65  
 calculations 114–115  
 transition metals 96–97  
 chemical properties 97  
 physical properties 96–97  
 transparent materials 202

## U

universal indicator 52  
 unsaturated compounds 161, 173

## V

valency 41  
 variables 136  
 viscosity 153  
 volumetric flasks 112

## W

water  
 distillation 10, 14  
 for chemical analysis 14  
 for drinking 14–15  
 weak acids 55

## X

X-rays 28

## Y

yields from reactions 108–109, 122



# The Periodic Table of the Elements

1	2	3	4	5	6	7	0
<div> <div>1</div> <div>H</div> <div>hydrogen</div> <div>1</div> </div>	Key						<div> <div>4</div> <div>He</div> <div>helium</div> <div>2</div> </div>
	<div> <div>relative atomic mass</div> <div>atomic symbol</div> <div>name</div> <div>atomic (proton) number</div> </div>						
<div> <div>7</div> <div>Li</div> <div>lithium</div> <div>3</div> </div>	<div> <div>9</div> <div>Be</div> <div>beryllium</div> <div>4</div> </div>						<div> <div>11</div> <div>B</div> <div>boron</div> <div>5</div> </div>
<div> <div>23</div> <div>Na</div> <div>sodium</div> <div>11</div> </div>	<div> <div>24</div> <div>Mg</div> <div>magnesium</div> <div>12</div> </div>						<div> <div>12</div> <div>C</div> <div>carbon</div> <div>6</div> </div>
		<div> <div>45</div> <div>Sc</div> <div>scandium</div> <div>21</div> </div>	<div> <div>48</div> <div>Ti</div> <div>titanium</div> <div>22</div> </div>	<div> <div>51</div> <div>V</div> <div>vanadium</div> <div>23</div> </div>	<div> <div>52</div> <div>Cr</div> <div>chromium</div> <div>24</div> </div>	<div> <div>55</div> <div>Mn</div> <div>manganese</div> <div>25</div> </div>	<div> <div>56</div> <div>Fe</div> <div>iron</div> <div>26</div> </div>
<div> <div>39</div> <div>K</div> <div>potassium</div> <div>19</div> </div>	<div> <div>40</div> <div>Ca</div> <div>calcium</div> <div>20</div> </div>	<div> <div>89</div> <div>Y</div> <div>yttrium</div> <div>39</div> </div>	<div> <div>91</div> <div>Zr</div> <div>zirconium</div> <div>40</div> </div>	<div> <div>93</div> <div>Nb</div> <div>niobium</div> <div>41</div> </div>	<div> <div>96</div> <div>Mo</div> <div>molybdenum</div> <div>42</div> </div>	<div> <div>[98]</div> <div>Tc</div> <div>technetium</div> <div>43</div> </div>	<div> <div>101</div> <div>Ru</div> <div>ruthenium</div> <div>44</div> </div>
<div> <div>85</div> <div>Rb</div> <div>rubidium</div> <div>37</div> </div>	<div> <div>88</div> <div>Sr</div> <div>strontium</div> <div>38</div> </div>						<div> <div>103</div> <div>Rh</div> <div>rhodium</div> <div>45</div> </div>
		<div> <div>139</div> <div>La*</div> <div>lanthanum</div> <div>57</div> </div>	<div> <div>178</div> <div>Hf</div> <div>hafnium</div> <div>72</div> </div>	<div> <div>181</div> <div>Ta</div> <div>tantalum</div> <div>73</div> </div>	<div> <div>184</div> <div>W</div> <div>tungsten</div> <div>74</div> </div>	<div> <div>186</div> <div>Re</div> <div>rhennium</div> <div>75</div> </div>	<div> <div>190</div> <div>Os</div> <div>osmium</div> <div>76</div> </div>
<div> <div>133</div> <div>Cs</div> <div>caesium</div> <div>55</div> </div>	<div> <div>137</div> <div>Ba</div> <div>barium</div> <div>56</div> </div>						<div> <div>192</div> <div>Ir</div> <div>iridium</div> <div>77</div> </div>
		<div> <div>[227]</div> <div>Ac*</div> <div>actinium</div> <div>89</div> </div>	<div> <div>[261]</div> <div>Rf</div> <div>rutherfordium</div> <div>104</div> </div>	<div> <div>[262]</div> <div>Db</div> <div>dundubium</div> <div>105</div> </div>	<div> <div>[266]</div> <div>Sg</div> <div>seaborgium</div> <div>106</div> </div>	<div> <div>[264]</div> <div>Bh</div> <div>bohrium</div> <div>107</div> </div>	<div> <div>[277]</div> <div>Hs</div> <div>hassium</div> <div>108</div> </div>
<div> <div>[223]</div> <div>Fr</div> <div>francium</div> <div>87</div> </div>	<div> <div>[226]</div> <div>Ra</div> <div>radium</div> <div>88</div> </div>						<div> <div>[268]</div> <div>Mt</div> <div>meitnerium</div> <div>109</div> </div>
							<div> <div>[271]</div> <div>Ds</div> <div>darmstadtium</div> <div>110</div> </div>
							<div> <div>[272]</div> <div>Rg</div> <div>roentgenium</div> <div>111</div> </div>
							<div> <div>201</div> <div>Hg</div> <div>mercury</div> <div>80</div> </div>
							<div> <div>204</div> <div>Tl</div> <div>thallium</div> <div>81</div> </div>
							<div> <div>207</div> <div>Pb</div> <div>lead</div> <div>82</div> </div>
							<div> <div>209</div> <div>Bi</div> <div>bismuth</div> <div>83</div> </div>
							<div> <div>[209]</div> <div>Po</div> <div>polonium</div> <div>84</div> </div>
							<div> <div>[210]</div> <div>At</div> <div>astatine</div> <div>85</div> </div>
							<div> <div>[222]</div> <div>Rn</div> <div>radon</div> <div>86</div> </div>
							<div> <div>127</div> <div>I</div> <div>iodine</div> <div>53</div> </div>
							<div> <div>131</div> <div>Xe</div> <div>xenon</div> <div>54</div> </div>
							<div> <div>128</div> <div>Te</div> <div>tellurium</div> <div>52</div> </div>
							<div> <div>75</div> <div>As</div> <div>arsenic</div> <div>33</div> </div>
							<div> <div>79</div> <div>Se</div> <div>selenium</div> <div>34</div> </div>
							<div> <div>80</div> <div>Br</div> <div>bromine</div> <div>35</div> </div>
							<div> <div>84</div> <div>Kr</div> <div>krypton</div> <div>36</div> </div>
							<div> <div>70</div> <div>Ga</div> <div>gallium</div> <div>31</div> </div>
							<div> <div>73</div> <div>Ge</div> <div>germanium</div> <div>32</div> </div>
							<div> <div>115</div> <div>In</div> <div>indium</div> <div>49</div> </div>
							<div> <div>119</div> <div>Sn</div> <div>tin</div> <div>50</div> </div>
							<div> <div>112</div> <div>Cd</div> <div>cadmium</div> <div>48</div> </div>
							<div> <div>63.5</div> <div>Cu</div> <div>copper</div> <div>29</div> </div>
							<div> <div>65</div> <div>Zn</div> <div>zinc</div> <div>30</div> </div>
							<div> <div>59</div> <div>Co</div> <div>cobalt</div> <div>27</div> </div>
							<div> <div>59</div> <div>Ni</div> <div>nickel</div> <div>28</div> </div>
							<div> <div>27</div> <div>Al</div> <div>aluminium</div> <div>13</div> </div>
							<div> <div>31</div> <div>P</div> <div>phosphorus</div> <div>15</div> </div>
							<div> <div>32</div> <div>S</div> <div>sulfur</div> <div>16</div> </div>
							<div> <div>35.5</div> <div>Cl</div> <div>chlorine</div> <div>17</div> </div>
							<div> <div>19</div> <div>F</div> <div>fluorine</div> <div>9</div> </div>
							<div> <div>16</div> <div>O</div> <div>oxygen</div> <div>8</div> </div>
							<div> <div>14</div> <div>N</div> <div>nitrogen</div> <div>7</div> </div>
							<div> <div>20</div> <div>Ne</div> <div>neon</div> <div>10</div> </div>
							<div> <div>40</div> <div>Ar</div> <div>argon</div> <div>18</div> </div>
							<div> <div>207</div> <div>Pb</div> <div>lead</div> <div>82</div> </div>
							<div> <div>209</div> <div>Bi</div> <div>bismuth</div> <div>83</div> </div>
							<div> <div>[209]</div> <div>Po</div> <div>polonium</div> <div>84</div> </div>
							<div> <div>[210]</div> <div>At</div> <div>astatine</div> <div>85</div> </div>
							<div> <div>[222]</div> <div>Rn</div> <div>radon</div> <div>86</div> </div>

\*The lanthanoids (atomic numbers 58-71) and the actinoids (atomic numbers 90-103) have been omitted.

The relative atomic masses of copper and chlorine have not been rounded to the nearest whole number.

Published by Pearson Education Limited, 80 Strand, London, WC2R 0RL.

[www.pearsonschoolsandcolleges.co.uk](http://www.pearsonschoolsandcolleges.co.uk)

Copies of official specifications for all Edexcel qualifications may be found on the website: [www.edexcel.com](http://www.edexcel.com)

Text © Mark Levesley, Iain Brand, Nigel Saunders, Sue Robilliard, John Ling, Penny Johnson and Pearson Education Ltd 2016

Series editor: Mark Levesley

Designed by Poppy Marks, Pearson Education Limited

Typeset by Phoenix Photosetting, Chatham, Kent

Original illustrations © Pearson Education Limited 2016

Illustrated by KJA Artists Illustration Agency and Phoenix Photosetting, Chatham, Kent

Cover design by Poppy Marks and Colin Tilley Loughrey

Picture research by Rebecca Sodergren

Cover photo © 123RF.com: Aafak Cakar, Maksim Kostenko;

SuperStock: Corbis

The rights of Mark Levesley, Iain Brand, Nigel Saunders, Sue Robilliard and John Ling to be identified as authors of this work have been asserted by them in accordance with the Copyright, Designs and Patents Act 1988.

First published 2016

19 18 17 16

10 9 8 7 6 5 4 3 2 1

#### British Library Cataloguing in Publication Data

A catalogue record for this book is available from the British Library

ISBN 9781292120218

#### Copyright notice

All rights reserved. No part of this publication may be reproduced in any form or by any means (including photocopying or storing it in any medium by electronic means and whether or not transiently or incidentally to some other use of this publication) without the written permission of the copyright owner, except in accordance with the provisions of the Copyright, Designs and Patents Act 1988 or under the terms of a licence issued by the Copyright Licensing Agency, Saffron House, 6–10 Kirby Street, London EC1N 8TS ([www.cla.co.uk](http://www.cla.co.uk)). Applications for the copyright owner's written permission should be addressed to the publisher.

Printed in Slovakia by Neografia.

#### A note from the publisher

In order to ensure that this resource offers high-quality

support for the associated Pearson qualification, it has been through a review process by the awarding body. This process confirms that this resource fully covers the teaching and learning content of the specification or part of a specification at which it is aimed. It also confirms that it demonstrates an appropriate balance between the development of subject skills, knowledge and understanding, in addition to preparation for assessment.

Endorsement does not cover any guidance on assessment activities or processes (e.g. practice questions or advice on how to answer assessment questions), included in the resource nor does it prescribe any particular approach to the teaching or delivery of a related course.

While the publishers have made every attempt to ensure that advice on the qualification and its assessment is accurate, the official specification and associated assessment guidance materials are the only authoritative source of information and should always be referred to for definitive guidance.

Pearson examiners have not contributed to any sections in this resource relevant to examination papers for which they have responsibility. Examiners will not use endorsed resources as a source of material for any assessment set by Pearson.

Endorsement of a resource does not mean that the resource is required to achieve this Pearson qualification, nor does it mean that it is the only suitable material available to support the qualification, and any resource lists produced by the awarding body shall include this and other appropriate resources.

#### Acknowledgements

The following authors contributed text to previous Pearson publications and the publishers are grateful for their permission to include elements of their work: Richard Grime, Ray Oliver, Gemma Young, Penny Johnson and Peter Ellis. The publishers would like to thank Steve Gray and Penny Johnson for their original contributions.

The author and publisher would like to thank the following individuals and organisations for permission to reproduce photographs, figures and text:

#### Photographs

(Key: b-bottom; c-centre; l-left; r-right; t-top)

SC1-2 TheOceanCleanup.com; SC1a Alamy Images: Arctic Images (A); SC2a Alamy Images: studiomode (C). Pearson Education Ltd: (B). Shutterstock.com: sunny337 (A); SC2b Alamy Images: Vicki Beaver (A). Science Photo Library Ltd: Mark de Fraeye (C); Javier Trueba / MSF (B); SC2c Corbis: Anthony Rakusen / Cultura (B). Getty Images: Francois Guillot / AFP (A); SC2d Fotolia.com: nikkytok (A); SC2d CP



Shutterstock.com: Ralko (A); SC2e Pearson Education Ltd: (C).  
Shutterstock.com: Timothy Epp (A)

SC3 Reproduced with permission of D. Pearmain, S. J. Park, Z. W. Wang, A. Abdela, R. E. Palmer and Z. Y. Li; University of Birmingham; SC3a Getty Images: Tom Shaw (D). Pearson Education Ltd: Tsz-shan Kwok (A); SC3b Shutterstock.com: Paul Fleet (C); SC3c Getty Images: Corbis / Nagasaki Atomic Bomb Museum / epa (B)

SC4 Science Photo Library Ltd; SC4a NikNaks: (B).  
Shutterstock.com: Baloncici (A); SC4b Alamy Images: Jon Bower UK (A); SC4c Shutterstock.com: Domenic Gareri (A)

SC5-7 University of California, Berkeley: Felix R. Fischer Ph.D, Department of Chemistry; SC5a Science Photo Library Ltd: Andrew Lambert Photography (A); SC5b Science Photo Library Ltd: Gary Hincks (C); Dirk Wiersma (B); SC5c Science Photo Library Ltd: Tony Camacho (B); SC7a Getty Images: Photolibrary / Huw Jones (B). Science Photo Library Ltd: Alvis Uptis / AgstockUSA (C); SC7b Shutterstock.com: ribeiroantonio (D); SC7c Fotolia.com: Silvano Rebai (D). Getty Images: Image Ideas (A); SC7d Science Photo Library Ltd: Indigo Molecular Images (A)

SC8 Shutterstock.com: Science Photo Library; SC8a Martyn F. Chillmaid: (A, B); SC8b Martyn F. Chillmaid: (F). Pearson Education Ltd: Trevor Clifford (C). Science Photo Library Ltd: GIPhotoStock (A); SC8c Shutterstock.com: Industry and Travel (A); SC8c CP Science Photo Library Ltd: Andrew Lambert Photography (C); SC8d Science Photo Library Ltd: Mid Essex Hospital Services NHS Trust (A); Martyn F. Chillmaid (B); SC8d CP Science Photo Library Ltd: Trevor Clifford Photography (A); GIPhotoStock (B); SC8e Martyn F. Chillmaid: (C). Science Photo Library Ltd: Martyn F. Chillmaid (B). Shutterstock.com: Paul Prescott (A); SC8f Getty Images: Popperfoto (A). Martyn F. Chillmaid: (C). Pearson Education Ltd: Trevor Clifford (B); SC8g Science Photo Library Ltd: ER Degginger (A)

SC9 Getty Images: Corbis / Caroline; SC9a Getty Images: Auscape / UIG (C); SC9b Fundamental Photographs: Richard Megna (B). Science Photo Library Ltd: Martyn F. Chillmaid (A); SC9c Science Photo Library Ltd: Andrew Lambert Photography (A) SC10-11 Alamy Images: age footstock; SC10a Alamy Images: WENN UK (C); SC11a Science Photo Library Ltd: Charles D. Winters (B). The Open University: (A); SC11b Dr. Antony van der Ent: Sustainable Minerals Institute / The University of Queensland (C). Shutterstock.com: Steffen Foerster (A); SC11c Courtesy of the Jersey Heritage Collection: (C). Shutterstock.com: Ron Ellis (D); icarmen13 (A); SC11d Alamy Images: Stephen Saks Photography (B); AF archive (D). Ocean Sole <http://www.ocean-sole.com/>: (A); SC12a Fundamental Photographs: Chip Clark (D). Pearson Education Ltd: Trevor Clifford (A); SC12b Alamy Images: Corbin17 (A). Science Photo Library Ltd: Frans Lanting,

Mint Images (B); SC13a Alamy Images: sciencephotos (D). Shutterstock.com: Tinalmages (E); SasinT (A); SC13b Fotolia.com: pieropoma (D). Pearson Education Ltd: Gareth Boden (B). Shutterstock.com: S.Borisov (A); AHPix (C); SC13c Fotolia.com: Ilya Akinshin (A). Shutterstock.com: optimarc (D); Madlen (B); SC13d Alamy Images: David Hunter (A). Fotolia.com: zhu difeng (B). Shutterstock.com: Vereshchagin Dmitry (C); SC13e Alamy Images: TravelCollection (B); Heather Drake (C). Fotolia.com: kamilpetran (A); Andrey Armyagov (D)

SC14-16 Shutterstock.com: Jason.Lee; SC14a Alamy Images: robertharding (C); Branislav Bokun (B). Getty Images: Christian Science Monitor (A); SC14b Alamy Images: studiomode (B). Edinburgh Napier University: (C); SC14c Science Photo Library Ltd: Martyn F. Chillmaid (A, C); SC14d CP Science Photo Library Ltd: Martyn F. Chillmaid (A, B, C); SC14e Getty Images: Roger Viollet (A). Pearson Education Ltd: Trevor Clifford (C); SC15a Science Photo Library Ltd: US Department of Agriculture (A); SC15b Science Photo Library Ltd: Keith Kent (A); SC16a Alamy Images: Jim West (A). Science Photo Library Ltd: (B)

SC17-19 Shutterstock.com: PawelG Photo; SC17a Pearson Education Ltd: Trevor Clifford (B); SC17b Science Photo Library Ltd: Andrew Lambert Photography (A, C); SC17c Science Photo Library Ltd: Martyn F. Chillmaid (B); Andrew Lambert Photography (C); SC17d Getty Images: Nick Obank / Barcroft Media (D). Shutterstock.com: Karin Hildebrand Lau (E); SC18a Alamy Images: Bon Appetit (A); SC18b Getty Images: (B); SC18b CP Science Photo Library Ltd: Martyn F. Chillmaid (B); SC18c Science Photo Library Ltd: (A); SC19a Science Photo Library Ltd: (A). Shutterstock.com: M. Norris (B); SC19b Shutterstock.com: Stefan Holm (A)

SC20-21 Alamy Images: Plane Mad; SC20a Alamy Images: Daniel J. Cox (A). Getty Images: loonger (C). Shutterstock.com: Dmitry Naumov (D); SC20b Science Photo Library Ltd: Hybrid Images (B). Shutterstock.com: jocic (A); SC20c Alamy Images: Klaus Oskar Bromberg (A); SC20d Alamy Images: Alan Keith Beastall (D). Pearson Education Ltd: Trevor Clifford (B). Science Photo Library Ltd: (A). Shutterstock.com: Danny E Hooks (C); SC20e Alamy Images: Wayne Hutchinson (C). Science Photo Library Ltd: Simon Fraser (A); Monica Schroeder (B). Shutterstock.com: Hung Chung Chih (D); SC20f Reuters: Alex Gallardo (D). Shutterstock.com: NicVW (A); SC21a Alamy Images: World History Archive (A). Science Photo Library Ltd: Natural History Museum, London (D); NASA / JPL (B); SC21b Science Photo Library Ltd: Martyn F. Chillmaid (D); SC21c Alamy Images: Arctic Images (D); SC21d 123RF.com: warawoot nanta (A). Science Photo Library Ltd: Tony Camacho (B); Georgette Douwma (C)

SC22- 24 Alamy Images: epa european pressphoto agency b.v.; SC22a Alamy Images: Clarence Holmes Wildlife (A); SC22b NASA: (A). Pearson Education Ltd: Trevor Clifford

(C). **Shutterstock.com**: Vlad Ageshin (B); **SC23a Fotolia.com**: ra3rn (A/vodka); Givaga (A/whisky). **Science Photo Library Ltd**: Martyn F. Chillmaid (C); (D). **Shutterstock.com**: Nitr (A/beer); Dmitri Gristenko (A/wine); **SC23b CP Alamy Images**: inga spence (A); **SC23c Alamy Images**: Mediablitzimages (B). **Martyn F. Chillmaid**: (D); **SC24a Artville**: Dennis Nolan (A). **PhotoDisc**: (D); **SC24b 123RF.com**: (A/bag, A/pan). **Shutterstock.com**: pryzmat (A/pipes); paijicha (A/bottle top); CrackerClips Stock Media (A/packing chips); **SC24c Alamy Images**: Alex Segre (B). **Science Photo Library Ltd**: Trevor Clifford (A); **SC24d Alamy Images**: Green Stock Media (D). **Getty Images**: Mint Images (B). **Science Photo Library Ltd**: Simon Fraser (A)

**SC25-26 Science Photo Library Ltd**: Patrick Landmann; **SC25a Alamy Images**: Roy Conchie (A). **Science Photo Library Ltd**: (B); **SC25b Pearson Education Ltd**: Trevor Clifford (C). **Science Photo Library Ltd**: Martyn F. Chillmaid (B, D). **Shutterstock.com**: K Steve Cope (A); **SC25c Alamy Images**: David Chapman (A). **Pearson Education Ltd**: Trevor Clifford (D). **Science Photo Library Ltd**: Trevor Clifford Photography (B); Charles D. Winters (C); **SC25c CP Shutterstock.com**: anyaivanova (A); **SC26a Alamy Images**: ITAR-TASS Photo Agency (C); Apelöga AB (B). **Fotolia.com**: slavun (A); ronstik (D); **SC26b Bart van Overbeeke Fotografie**: (A). **Getty Images**: matsou (B); **SC26c Science Photo Library Ltd**: Royal Institution Of Great Britain (B); Eye of Science (C); (A)

Cover images: Front: **123RF.com**: Aafak Cakar, Maksim Kostenko; **SuperStock**: Corbis

All other images © Pearson Education

### Text

SC3 Atomic Structure opener paragraph on page 17 adapted from Professor Richard Palmer, The Nanoscale Physics Research Laboratory, University of Birmingham. Reproduced with the permission of Professor Richard Palmer.

### Websites

Pearson Education Limited is not responsible for the content of any external internet sites. It is essential for tutors to preview each website before using it in class so as to ensure that the URL is still accurate, relevant and appropriate. We suggest that tutors bookmark useful websites and consider enabling students to access them through the school/college intranet.